

Query Match 100.0%; Score 32; DB 6; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Dy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 4
ACD99725
ID ACD99725 standard; DNA; 32 BP.
XX
AC ACD99725;
XX
DT 25-SEP-2003 (first entry)
XX
DE Immunostimulatory nucleic acid #411.
XX
KW Immunostimulatory; antiinflammatory; dermatological; antipruritic;
KW antilucer; gene therapy; vaccine; non-allergic inflammatory disease;
KW psoriasis; eczema; allergic contact dermatitis; latex dermatitis;
KW inflammatory bowel disease; ulcerative colitis; Crohn's disease; ss.
XX
OS Synthetic.
XX
PN US2003050268-A1.
XX
PD 13-MAR-2003.
XX
PF 29-MAR-2002; 2002US-00112653.
XX
PR 29-MAR-2001; 2001US-0279642P.
XX
PA (KRIE/) KRIEG A M.
PA (BERG/) BERG D J.
XX
PI Krieg AM, Berg DJ;
XX
DR WPI; 2003-521815/49.
XX
PT Treating non-allergic inflammatory diseases, such as psoriasis, eczema,
PT allergic contact dermatitis, latex dermatitis or inflammatory bowel
PT disease by administering an immunostimulatory nucleic acid.
XX
PS Disclosure; Page 19; 229pp; English.
XX
CC The invention describes a method of treating non-allergic inflammatory
CC disease comprising administering to a subject having or at risk of
CC developing a non-allergic inflammatory disease an immunostimulatory
CC nucleic acid for prevention or treatment of the disease. The method is
CC useful for treating non-allergic inflammatory diseases, such as
CC psoriasis, eczema, allergic contact dermatitis, latex dermatitis or
CC inflammatory bowel disease e.g., ulcerative colitis or Crohn's disease.
CC This sequence represents an immunostimulatory nucleic acid
XX
SQ Sequence 32 BP; 0 A; 5 C; 8 G; 19 T; 0 U; 0 Other;

Query Match 100.0%; Score 32; DB 9; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Dy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 5
ADB36802
ID ADB36802 standard; DNA; 32 BP.
XX
AC ADB36802;

XX
DT 04-DEC-2003 (first entry)
XX
DE Immunostimulatory nucleic acid #416.
XX
KW de; allergy; asthma; poly-G nucleic acid; aerosol formulation;
KW hypo-responsive subject; immunostimulatory.
XX
OS Synthetic.
XX
PN US2003087848-A1.
XX
PD 08-MAY-2003.
XX
PF 02-FEB-2001; 2001US-00776479.
XX
PR 03-FEB-2000; 2000US-0179991P.
XX
PA (BRAT/) BRATZLER R L.
PA (PETER/) PETERSEN D M.
PA (FOUR/) FOURON Y.
XX
PI Bratzler RL, Petersen DM, Fouron Y;
XX
DR WPI; 2003-657977/62.
XX
PT Treating and/or preventing allergy or asthma using an immunostimulatory
PT nucleic acid alone or in combination with an asthma/allergy medicament.
XX
PS Disclosure; Page 11; 221pp; English.
XX
CC The invention relates to a method of treating or preventing allergy or
CC asthma which comprises administering to a subject a poly-G nucleic acid
CC in an aerosol formulation. The methods and compositions of the present
CC invention are useful for diagnosing and/or treating asthma and allergy
CC especially in a hypo-responsive subject. The present sequence represents
CC an immunostimulatory nucleic acid of the invention.
XX
SQ Sequence 32 BP; 0 A; 5 C; 8 G; 19 T; 0 U; 0 Other;

Query Match 100.0%; Score 32; DB 9; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Dy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 6
ADD89955
ID ADD89955 standard; DNA; 32 BP.
XX
AC ADD89955;
XX
DT 29-JAN-2004 (first entry)
XX
DE CPG oligonucleotide used in immunostimulant complex.
XX
KW Immunostimulant; vaccine; CPG; adjuvant; ss.
XX
OS Synthetic.
XX
PN WO2003068169-A2.
XX
PD 21-AUG-2003.
XX
PF 14-FEB-2003; 2003WO-US004711.
XX
PR 14-FEB-2002; 2002US-00076674.
PR 31-JAN-2003; 2003US-00076674.
XX
PA (UNBI-) UNITED BIOMEDICAL INC.

XX Sokol1 KK;
 PI
 XX
 DR WPI; 2003-778890/73.
 XX
 PT Stabilized immunostimulating complex, useful for vaccination, e.g.
 XX against human immune deficiency viruses, comprises cationic peptide
 PS immunogen and anionic oligonucleotide.
 XX
 XX Claim 10; SEQ ID NO 1, 159pp; English.
 CC The present sequence is that of an Cpg oligonucleotide, denoted Cpg1,
 CC that is used in immunostimulatory complexes specifically adapted to act
 CC as adjuvant and as peptide immunogen stabiliser. The complexes comprise a
 CC Cpg oligonucleotide and a biologically active peptide immunogen. The
 CC complex is particularly and efficiently present peptide immunogen. The
 CC cells of the immune system to produce an immune response. The
 CC complexes may be prepared with various ratios of peptides to Cpg
 CC oligonucleotides to provide different physical properties, such as the
 CC size of the microparticle. Cpg1 possesses a Cpg motif sequestered within
 CC in vivo and improved stability by binding stronger adjuvantation
 CC affinites than shorter oligonucleotides. Peptide immunogens used in
 CC claimed immunostimulatory complexes are: LHRH peptides useful in a
 CC vaccine for prostate cancer immunotherapy; CD4 peptides useful in an anti
 CC -CD4 immunotherapeutic vaccine for the treatment of HIV infection; IgS
 CC peptides useful in an allergy vaccine; and foot-and-mouth-disease
 XX
 SQ Sequence 32 BP; 0 A; 5 C; 8 G; 19 T; 0 U; 0 Other;
 Query Match 100.0%; Score 32; DB 10; Length 32;
 Best Local Similarity 100.0%; Pred. No. 1.1e-07;
 Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
 DB 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 7
 AD56900
 ID AD56900 standard; DNA; 32 BP.
 XX
 AC AD56900;
 XX
 DT 06-MAY-2004 (first entry)
 XX
 DE Cpg oligonucleotide, Cpg1.
 XX
 KW Immunostimulatory complex; adjuvant; peptide immunogen stabiliser;
 KW water-in-oil emulsion; suspension; vaccine; prostate cancer;
 KM hormone ablation; allergy; HIV infection; foot-and-mouth disease;
 XX therapy; ss.
 OS Synthetic.
 XX
 PN US2004009897-A1.
 XX
 PD 15-JAN-2004.
 XX
 PF 21-MAY-2003; 2003US-00355161.
 XX
 PR 14-FEB-2002; 2002US-00076674.
 XX
 PA (SOKO/) SOKOL K K.
 XX
 PI Sokol1 KK;
 XX
 DR WPI; 2004-212745/20.
 XX
 PT Stabilized immunostimulatory complex useful for treating allergy, HIV
 infection or prostate cancer, comprising cationic peptide immunogen and

PT anionic Cpg oligonucleotide.
 XX
 XX Claim 10; SEQ ID NO 1; 63pp; English.
 PS
 CC The invention relates to an immunostimulatory complex specifically
 CC adapted to act as adjuvant and as a peptide immunogen stabiliser. The
 CC invention is useful for preparing a water-in-oil emulsion, suspension and
 CC vaccine. It is also useful for treating prostate cancer, hormone
 CC ablation, allergy, HIV infection, foot-and-mouth disease, etc. The
 CC present sequence is an oligonucleotide used in the invention.
 XX
 SQ Sequence 32 BP; 0 A; 5 C; 8 G; 19 T; 0 U; 0 Other;
 Query Match 100.0%; Score 32; DB 12; Length 32;
 Best Local Similarity 100.0%; Pred. No. 1.1e-07;
 Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
 DB 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 8
 ACD91394
 ID ACD91394 standard; DNA; 27 BP.
 XX
 AC ACD91394;
 XX
 DT 22-SEP-2003 (first entry)
 XX
 DE Adjuvant-type Cpg containing oligonucleotide #3.
 XX
 DE Cpg island; ss; HIV infection; gene therapy; vaccine; B-cell;
 KW immunostimulatory; adjuvant.
 KM
 XX
 OS Synthetic.
 XX
 PN US2003050263-A1.
 XX
 PD 13-MAR-2003.
 XX
 PF 16-AUG-2001; 2001US-00931583.
 XX
 PR 15-JUL-1994; 94US-00276358.
 PR 07-FEB-1995; 95US-00386063.
 PR 08-OCT-1999; 99US-00415142.
 XX
 PA (IOWA) UNIV IOWA RES FOUND.
 XX
 PI Krieg AM, Kliman D, Steinberg AD;
 XX
 DR WPI; 2003-512356/48.
 XX
 PT Treating a subject infected with HIV by administering a Cpg nucleic acid.
 XX
 PS Claim 50; Page 20; 22pp; English.
 XX
 CC The invention relates to treating a subject infected with HIV comprising
 CC administering a Cpg nucleic acid (e.g. an adjuvant type Cpg
 CC oligonucleotide, an immunostimulatory Cpg oligonucleotide or a B cell
 CC stimulatory Cpg oligonucleotide). The Cpg are used as gene therapy
 CC vaccines to treat a subject infected with HIV. The present sequence is an
 CC adjuvant type Cpg oligonucleotide
 XX
 SQ Sequence 27 BP; 0 A; 4 C; 6 G; 17 T; 0 U; 0 Other;
 Query Match 81.2%; Score 26; DB 9; Length 27;
 Best Local Similarity 100.0%; Pred. No. 0.00016;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 TCGTCGTTTGTGCGTTTGTGCGTTT 26
 DB 1 TCGTCGTTTGTGCGTTTGTGCGTTT 26


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RESULT 9
ID AAF99178 standard; DNA; 29 BP.
AC AAF99178;
XX
DT 12-JUN-2001 (first entry)
DE Immunostimulatory nucleic acid #294.
KM Vaccine; cytotoxic; virucidal; bactericidal; fungicidal; anti-parasitic;
KW immunostimulatory; tumour; viral infection; bacterial infection;
KV fungal infection; parasitic infection; cancer; asthma;
KX infectious disease; allergy; immune deficiency; phosphorochioate; ss.
OS Synthetic.
PN WO200122972-A2.
PD 05-APR-2001.
PE 25-SEP-2000; 2000MO-USO26383.
PR 25-SEP-1999; 99US-0156113P.
PR 27-SEP-1999; 99US-0156135P.
PR 23-AUG-2000; 2000US-0227436P.
PA (IOWA ) UNIV IOWA RES FOUND.
PA (COLE-) COLEY PHARM GMBH.
PL Kriegl AM, Schetter C, Vollmer J;
XX WPI; 2001-273485/28.
PT Vaccinating against tumors, infectious diseases, allergies and asthma
PT using immunostimulatory Py-rich and TG nucleic acids.
PS Claim 101; Page 44; 338pp; English.
XX
XX The present invention relates to a method for stimulating an immune
XX response. The method comprises administering an immunostimulatory nucleic
XX acid to a non-rodent subject in sufficient quantity to stimulate an
XX immune response. The present sequence is one such immunostimulatory
XX nucleic acid. The immunostimulatory nucleic acids can be pyrimidine rich
XX (py-rich) or thymidine (T) rich. The method is used to vaccinate subjects
XX against tumour antigens, viral antigens (e.g. herpesviridae, retroviridae
XX and/or toxomoviridae), bacterial antigens (e.g. toxoplasma,
XX haemophilus, campylobacter, clostridium, Escherichia coli and/or
XX staphylococcus), fungal antigens and/or parasitic antigens. The method is
XX also useful for preventing cancer, asthma, infectious disease, allergy or
XX immune deficiency. The present sequence can also be used to redirect a
XX Th2 to a Th1 immune response and to activate immune cells. Note: the
XX present sequence may have a phosphorochioate backbone
XX
SQ Sequence 29 BP; 0 A; 4 C; 6 G; 19 T; 0 U; 0 Other;
OY Query Match 81.2%; Score 26; DB 4; Length 29;
ID Best Local Similarity 100.0%; Pred.No. 0.00016;
XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX 1 TCATCGTTTGTCGTTTGTCGTTT 26
XX 1 TCATCGTTTGTCGTTTGTCGTTT 26
XX
RESULT 10
ID ABS77821 standard; DNA; 29 BP.
AC ABS77821;
XX
XX ABS77821;
XX
```

DT	13-DEC-2002	(first entry)	
XX		Angiogenesis inhibitory oligonucleotide #305.	
DE		Angiogenesis inhibitor; ss; angiogenesis; solid tumour growth;	
KW		tumour metastasis; precancerous lesion; rheumatoid arthritis; psoriasis;	
KW		diabetic retinopathy; retinopathy of prematurity; macular degeneration;	
KW		corneal graft rejection; neovascular glaucoma; retrolental fibroplasia;	
KW		rubeosis; Osler-Weber Syndrome; myocardial angiogenesis;	
KW		plaque neovascularisation; telangiectasia; haemophiliac joint;	
KW		angiofibroma; wound granulation; intestinal adhesion; atherosclerosis;	
KW		scleroderma; hypertrophic scar.	
XX			
OS		Synthetic.	
XX			
PN	WO200253141-A2.		
XX			
PD	11-JUL-2002.		
XX			
PF	14-DEC-2001; 2001WO-US048458.		
XX			
PR	14-DEC-2000; 2000US-0255534P.		
XX			
PA	(COLE-) COLEY PHARM GROUP INC.		
XX			
P1	Bratzler RL;		
XX			
DR	WPI; 2002-566690/60.		
XX			
PT	Inhibiting angiogenesis in a subject, involves administering at least one		
XX	antiangiogenic nucleic acid molecule to the subject.		
XX			
PS	Claim 2; Page 25; 276pp; English.		
XX			
CC	The invention relates to inhibiting angiogenesis in a subject, comprising		
CC	administering at least one antiangiogenic nucleic acid molecule. Also		
CC	included is a kit comprising a first container housing the antiangiogenic		
CC	nucleic acids, and instructions for administering them to a subject		
CC	having a condition characterised by unwanted angiogenesis. The method is		
CC	useful for inhibiting angiogenesis associated with solid tumour growth,		
CC	tumour metastasis, precancerous lesion, rheumatoid arthritis, psoriasis,		
CC	diabetic retinopathy, retinopathy of prematurity, macular degeneration,		
CC	corneal graft rejection, neovascular glaucoma, retrolental fibroplasia,		
CC	rubeosis, Osler-Weber Syndrome, myocardial angiogenesis, plaque		
CC	neovascularisation, telangiectasia, haemophiliac joints, angiofibroma,		
CC	wound granulation, intestinal adhesions, atherosclerosis, scleroderma and		
CC	hypertrophic scars. The present sequence is an antiangiogenic nucleic		
CC	acid of the invention		
XX			
XX			
SQ	Sequence 29 BP; 0 A; 4 C; 6 G; 19 T; 0 U; 0 Other;		
XX			
Query March	81.2%; Score 26; DB 6; Length 29;		
Best Local Similarity	100.0%; Pred. No. 0.00016;		
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;			
QY	1 TCGTCGTTTGTGCGTTTGTGCGTTT 26		
DB	1 TCGTCGTTTGTGCGTTTGTGCGTTT 26		
RESULT 11			
ABL38761			
ID	ABL38761 standard; DNA; 29 BP.		
AC	ABL38761;		
XX			
XX	16-APR-2002 (first entry)		
DE	Immunostimulatory nucleic acid SEQ ID NO: 131.		
KW	Antibody-induced cell lysis; cancer; immunostimulatory; CD20;		
XX	angiogenesis; metastasis; cytostatic; ss.		
XX			

OS Synthetic.
 XX Key Location/Qualifiers
 FH modified_base 1
 FT /*tag= a
 FT /mod_base= OTHER
 FT /note= "5' biotinylated"
 XX
 XX W0200197843-A2.
 XX
 XX PD 27-DEC-2001.
 XX
 XX PF 22-JUN-2001; 2001WO-US020154.
 XX
 XX PR 22-JUN-2000; 2000US-0213346P.
 XX
 XX PA (IOWA) UNIV IOWA RES FOUND.
 XX
 XX PI Weiner G, Hartmann G;
 XX
 XX DR WPI; 2002-154611/20.
 XX
 XX PT Treating or preventing cancer, such as basal cell carcinoma, comprises
 PT administering immunostimulatory nucleic acids that induce expression of
 PT cell surface antigens and antibodies to a subject having or at risk of
 PT developing cancer.
 XX
 XX PS Disclosure; Page 129; 312pp; English.
 XX
 XX CC The present invention relates to methods for treating or preventing
 CC cancer, involving administering to a subject having or at risk of
 CC developing cancer immunostimulatory nucleic acids that induce expression
 CC of cell surface antigens and antibodies. The methods are useful for
 CC treating or preventing cancer such as basal cell carcinoma, bladder
 CC cancer, bone cancer, brain and central nervous system (CNS) cancer,
 CC breast cancer, cervical cancer, colon and rectum cancer, connective
 CC tissue cancer, esophageal cancer, eye cancer, kidney cancer, larynx
 CC cancer, leukemia, liver cancer, lung cancer, Hodgkin's lymphoma, non-
 CC Hodgkin's lymphoma, melanoma, myeloma, oral cavity cancer, ovarian
 CC cancer, pancreatic cancer, prostate cancer, rhabdomyosarcoma, skin
 CC cancer, stomach cancer, testicular cancer, and uterine cancer. The
 CC present sequence is an immunostimulatory oligonucleotide described in the
 CC exemplification of the invention
 XX
 XX SQ Sequence 29 BP; 0 A; 4 C; 6 G; 19 T; 0 U; 0 Other;
 XX
 XX Query Match 81.2%; Score 26; DB 6; Length 29;
 XX Best Local Similarity 100.0%; Pred. No. 0.00016;
 XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 XX QY 1 TCGTCGTTTTCGTCGTTTTCGTCGTTT 26
 XX 1 TCGTCGTTTTCGTCGTTTTCGTCGTTT 26
 XX
 XX DB 1 TCGTCGTTTTCGTCGTTTTCGTCGTTT 26
 XX
 XX RESULT 12
 XX ACD99609
 XX ID ACD99609 standard; DNA; 29 BP.
 XX
 XX AC ACD99609;
 XX
 XX DT 25-SEP-2003 (first entry)
 XX
 XX DE Immunostimulatory nucleic acid #295.
 XX
 XX KW Immunostimulatory; antiinflammatory; dermatological; antipruritic;
 KW antiulcer; gene therapy; vaccine; non-allergic inflammatory disease;
 KW psoriasis; eczema; allergic contact dermatitis; latex dermatitis;
 XX inflammatory bowel disease; ulcerative colitis; Crohn's disease; ss.
 OS Synthetic.
 XX
 XX PN US2003050268-A1.

XX PD 13-MAR-2003.
 XX
 XX PF 29-MAR-2002; 2002US-00112653.
 XX
 XX PR 29-MAR-2001; 2001US-0279642P.
 XX
 XX PA (KRIE/) KRIEG A M.
 XX (BERG/) BERG D J.
 XX
 XX PI Krieg AM, Berg DJ;
 XX
 XX DR WPI; 2003-521815/49.
 XX
 XX PT Treating non-allergic inflammatory diseases, such as psoriasis, eczema,
 PT allergic contact dermatitis, latex dermatitis or inflammatory bowel
 PT disease by administering an immunostimulatory nucleic acid.
 XX
 XX PS Disclosure; Page 16; 229pp; English.
 XX
 XX CC The invention describes a method of treating non-allergic inflammatory
 CC disease comprising administering to a subject having or at risk of
 CC developing a non-allergic inflammatory disease an immunostimulatory
 CC nucleic acid for prevention or treatment of the disease. The method is
 CC useful for treating non-allergic inflammatory diseases, such as
 CC psoriasis, eczema, allergic contact dermatitis, latex dermatitis or
 CC inflammatory bowel disease e.g., ulcerative colitis or Crohn's disease.
 CC This sequence represents an immunostimulatory nucleic acid
 XX
 XX SQ Sequence 29 BP; 0 A; 4 C; 6 G; 19 T; 0 U; 0 Other;
 XX
 XX Query Match 81.2%; Score 26; DB 9; Length 29;
 XX Best Local Similarity 100.0%; Pred. No. 0.00016;
 XX Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 XX QY 1 TCGTCGTTTTCGTCGTTTTCGTCGTTT 26
 XX 1 TCGTCGTTTTCGTCGTTTTCGTCGTTT 26
 XX
 XX DB 1 TCGTCGTTTTCGTCGTTTTCGTCGTTT 26
 XX
 XX RESULT 13
 XX ADB36680
 XX ID ADB36680 standard; DNA; 29 BP.
 XX
 XX AC ADB36680;
 XX
 XX DT 04-DEC-2003 (first entry)
 XX
 XX DE Immunostimulatory nucleic acid #294.
 XX
 XX KW de; allergy; asthma; poly-G nucleic acid; aerosol formulation;
 KW hypo-responsive subject; immunostimulatory.
 OS Synthetic.
 XX
 XX PN US2003087848-A1.
 XX
 XX PD 08-MAY-2003.
 XX
 XX PF 02-FEB-2001; 2001US-00776479.
 XX
 XX PR 03-FEB-2000; 2000US-0179991P.
 XX
 XX PA (BRATZ/) BRATZLER R L.
 PA (PETERSEN/) PETERSEN D M.
 XX (FOUR/) FOURON Y.
 XX
 XX PI Bratzler RL, Petersen DM, Fouron Y;
 XX
 XX DR WPI; 2003-657977/62.
 XX
 XX PT Treating and/or preventing allergy or asthma using an immunostimulatory
 PT nucleic acid alone or in combination with an asthma/allergy medicament.

XX Disclousure; Page 9; 221pp; English.
 PS
 XX
 CC The invention relates to a method of treating or preventing allergy or
 CC asthma which comprises administering to a subject a poly-G nucleic acid
 CC in an aerosol formulation. The methods and compositions of the present
 CC invention are useful for diagnosing and/or treating asthma and allergy
 CC especially in a hypo-responsive subject. The present sequence represents
 CC an immunostimulatory nucleic acid of the invention.
 XX
 SQ Sequence 29 BP; 0 A; 4 C; 6 G; 19 T; 0 U; 0 Other;
 Query Match: 81.2%; Score 26; DB 9; Length 29;
 Best Local Similarity 100.0%; Pred. No. 0.00016;
 Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 1 TCGTCGTTTGTGCGTTTGTGCGTTT 26
 1 TCGTCGTTTGTGCGTTTGTGCGTTT 26
 Db
 RESULT 14
 ADE39672
 ID ADE39672 standard; DNA; 48 BP.
 XX
 AC ADE39672;
 XX
 DT 12-FEB-2004 (first entry)
 XX
 DE oligonucleotide ODN 7 (INX-2006) SEQ ID NO:7.
 XX
 KM cancer; vaccine; lipid-nucleic acid; LNA; tumour-associated antigen;
 KM Th-1 based immune response; cytostatic; gene therapy;
 KM tumour growth inhibition; tumour; human; ss.
 XX
 OS Synthetic.
 OS Homo sapiens.
 PH Key Location/Qualifiers
 FT modified_base 1..48
 FT /tag= a
 FT /mod_base= OTHER
 FT /note= "optionally phosphorothioate linkages and
 FT methylated cytosine residues"
 XX
 PN WO2003094828-A2.
 XX
 PD 20-NOV-2003.
 XX
 PF 12-MAY-2003; 2003WO-CA000679.
 XX
 PR 10-MAY-2002; 2002US-0379343P.
 PR 07-NOV-2002; 2002US-00290545.
 PR 04-APR-2003; 2003US-0460646P.
 XX
 PA (INEX-) INEX PHARM CORP.
 XX
 PI Tam YK, Sempke S, Klimuk S, Chikh G;
 XX
 DR WPI; 2004-011992/01.
 XX
 PT New cancer vaccine having a lipid-nucleic acid formulation in combination
 PT with at least one tumor-associated antigen, useful for stimulating
 PT enhanced responses against tumor-associated antigens and for inhibiting
 PT tumor growth.
 XX
 PS Example 9; SEQ ID NO 7; 119pp; English.
 XX
 CC The present invention describes a cancer vaccine (I), which comprises a
 CC lipid-nucleic acid (LNA) formulation in combination with at least one
 CC tumour-associated antigen that is mixed with or associated with the LNA
 CC formulation comprising a lipid component having at least one cationic
 CC lipid, and a nucleic acid component comprising at least one

CC oligonucleotide, where the vaccine is capable of stimulating a Th-1 based
 CC immune response in vivo to the at least one tumour-associated antigen.
 CC (I) has cytostatic activity, and can be used in vaccines, and in gene
 CC therapy. The methods and compositions of the present invention can be
 CC used for stimulating enhanced responses against tumour-associated
 CC antigens and for inhibiting tumour growth. The present sequence
 CC represents an oligonucleotide which is used in the exemplification of the
 CC present invention.
 XX
 SQ Sequence 48 BP; 0 A; 4 C; 12 G; 28 T; 0 U; 4 Other;
 Query Match: 78.1%; Score 25; DB 12; Length 48;
 Best Local Similarity 100.0%; Pred. No. 0.00051;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 1 TCGTCGTTTGTGCGTTTGTGCGTTT 25
 1 TCGTCGTTTGTGCGTTTGTGCGTTT 25
 Db
 RESULT 15
 AAV83725
 ID AAV83725 standard; DNA; 52 BP.
 XX
 AC AAV83725;
 XX
 DT 20-MAR-2003 (revised)
 DT 15-MAR-1999 (first entry)
 XX
 DE Plasmid pHIS Cpg-S motif containing oligonucleotide.
 XX
 KM Cpg-N motif; immunostimulation; antigen; Cpg-S motif; immunisation;
 KM viral antigen; bacterial antigen; parasite; therapeutic; growth factor;
 KM toxins; tumour suppressor; cytokine; apoptotic protein; interferon;
 KM hormone; clotting factor; ligand; receptor; ss.
 XX
 OS Synthetic.
 OS
 XX
 PN WO9852581-A1.
 XX
 PD 26-NOV-1998.
 XX
 PF 20-MAY-1998; 98WO-US010408.
 XX
 PR 20-MAY-1997; 97US-0047209P.
 PR 20-MAY-1997; 97US-0047233P.
 XX
 PA (OTTA-) OTTAMA CIVIC HOSPITAL LOEB RES INST.
 PA (IOMA) UNIV IOMA RES FOUND.
 PA (QIAG-) QIAGEN GMBH.
 XX
 PI Davis HL, Kriegl AM, Schorr J, Wu T;
 XX
 DR WPI; 1999-059712/05.
 XX
 PT Use of neutralising Cpg and stimulating Cpg motifs in DNA vectors - for
 PT enhancing the immunostimulatory effect of an antigen or enhancing the
 PT expression of a therapeutic polypeptide.
 XX
 PS Example 1; Page 41; 109pp; English.
 XX
 CC This sequence is used in the description of a method for enhancing the
 CC immunostimulatory effect of an antigen encoded by nucleic acid contained
 CC in a nucleic acid construct. The method involves determining the Cpg-N
 CC and Cpg-S motifs present in the construct, removing neutralising Cpg (Cpg
 CC -N) motifs and optionally inserting stimulatory Cpg (Cpg-S) motifs in the
 CC construct, thereby producing a nucleic acid construct having enhanced
 CC immunostimulatory efficacy. The method can be used for immunisation
 CC against viral antigens, e.g. from hepatitis B virus (HBV), bacterial
 CC antigens or an antigen derived from a parasite. They can also be used for
 CC expression of a therapeutic polypeptide, e.g. growth factors, toxins,
 CC tumour suppressors, cytokines, apoptotic proteins, interferons, hormones,
 CC clotting factors, ligands and receptors. (Updated on 20-MAR-2003 to

CC correct PA field.)
 XX
 SQ Sequence 52 BP; 1 A; 9 C; 14 G; 28 T; 0 U; 0 Other;
 Query Match 78.1%; Score 25; DB 2; Length 52;
 Best Local Similarity 100.0%; Pred. No. 0.0005;
 Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 TCGTCGTTTGTGTCGTTT 25
 Db 4 TCGTCGTTTGTGTCGTTT 28

Search completed: February 14, 2005, 06:52:40
 Job time : 298 secs

Query Match	100.0%	Score 32	DB 10	length 32
Best Local Similarity	100.0%	Pred. No. 1	1e-07	
Matches	32	Conservative	0	Mismatches 0; Indels 0; Gaps 0
QY	1	TCGTCGTTTGTGCGTTTGTGTCGTT	32	
Db	1	TCGTCGTTTGTGCGTTTGTGTCGTT	32	

RESULT 2
US-09-776-479-429
Sequence 429, Application US/09776479
Publication No. US20030087848A1
GENERAL INFORMATION:
APPLICANT: Bratzler, Robert L.
APPLICANT: Petersen, Deanna M.
TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
FILE REFERENCE: C1037/7013 (HCL/MAT)
CURRENT APPLICATION NUMBER: US/09/776, 479
PRIOR FILING DATE: 2001-02-02
NUMBER OF SEQ ID NOS: 1093
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 429
LENGTH: 32
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Sequence
US-09-776-479-429

Query Match
Best Local Similarity 100.0%; Score 32; DB 10; Length 32;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 3
US-09-776-479-429
Sequence 429, Application US/09776479
Publication No. US20040067902A9
GENERAL INFORMATION:
APPLICANT: Bratzler, Robert L.
APPLICANT: Petersen, Deanna M.
TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
FILE REFERENCE: C1037/7013 (HCL/MAT)
CURRENT APPLICATION NUMBER: US/09/776, 479
PRIOR FILING DATE: 2001-02-02
NUMBER OF SEQ ID NOS: 1093
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 429
LENGTH: 32
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Sequence
US-09-776-479-429

Query Match
Best Local Similarity 100.0%; Score 32; DB 11; Length 32;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 4
US-10-112-653-411
Sequence 411, Application US/10112653

Publication No. US20030050268A1
GENERAL INFORMATION:
APPLICANT: Krieg, Arthur M.
APPLICANT: Berg, Daniel J.
TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID FOR
FILE REFERENCE: C01039/70060 (AMS)
CURRENT APPLICATION NUMBER: US/10/112,653
PRIOR FILING DATE: 2002-03-29
NUMBER OF SEQ ID NOS: 1040
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 411
LENGTH: 32
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Oligonucleotide
US-10-112-653-411

Query Match
Best Local Similarity 100.0%; Score 32; DB 14; Length 32;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 5
US-10-017-995-429
Sequence 429, Application US/10017995
Publication No. US20030055014A1
GENERAL INFORMATION:
APPLICANT: Bratzler, Robert L.
TITLE OF INVENTION: Inhibition of Angiogenesis by Nucleic Acids
FILE REFERENCE: C1037/7025 (HCL/MAT)
CURRENT APPLICATION NUMBER: US/10/017, 995
PRIOR FILING DATE: 2001-12-18
NUMBER OF SEQ ID NOS: 1093
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 429
LENGTH: 32
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Sequence
US-10-017-995-429

Query Match
Best Local Similarity 100.0%; Score 32; DB 14; Length 32;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 6
US-10-076-674-1
Sequence 1, Application US/1007674
Publication No. US20030165478A1
GENERAL INFORMATION:
APPLICANT: Sokol, Kenneth K.
TITLE OF INVENTION: Stabilized Synthetic Immunogen Delivery System
FILE REFERENCE: C01039/70060 (AMS)
CURRENT APPLICATION NUMBER: US/10/076,674
PRIOR FILING DATE: 2002-04-23
NUMBER OF SEQ ID NOS: 11
SOFTWARE: PatentIn version 3.1

SEQ ID NO 1
LENGTH: 32
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: synthetic oligonucleotide
US-10-076-674-1

Query Match 100.0%; Score 32; DB 16; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 7

US-10-314-578-429
Sequence 429, Application US/10314578
Publication No. US20030212026A1
GENERAL INFORMATION:
APPLICANT: Krieger, Arthur M.
APPLICANT: Schetter, Christian
APPLICANT: Vollmer, Jorg
TITLE OF INVENTION: Immunostimulatory Nucleic Acids
FILE REFERENCE: C1039/7035 (HCL/MAT)
CURRENT APPLICATION NUMBER: US/10/314,578
CURRENT FILING DATE: 2002-12-09
PRIOR APPLICATION NUMBER: US 60/156,113
PRIOR FILING DATE: 1999-09-25
PRIOR APPLICATION NUMBER: US 60/156,135
PRIOR FILING DATE: 1999-09-27
PRIOR APPLICATION NUMBER: US 60/227,436
PRIOR FILING DATE: 2000-08-23
NUMBER OF SEQ ID NOS: 1145
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 429
LENGTH: 32
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Sequence
US-10-314-578-429

Query Match 100.0%; Score 32; DB 17; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

99

RESULT 8
US-10-355-161A-1
Sequence 1, Application US/10355161A
Publication No. US20040009897A1
GENERAL INFORMATION:
APPLICANT: Sokoll, Kenneth K.
TITLE OF INVENTION: Stabilized Synthetic Immunogen Delivery System
FILE REFERENCE: Immunogen Delivery System
CURRENT APPLICATION NUMBER: US/10/355,161A
CURRENT FILING DATE: 2003-01-31
PRIOR APPLICATION NUMBER: US 10/076674
PRIOR FILING DATE: 2002-02-14
NUMBER OF SEQ ID NOS: 13
SOFTWARE: PatentIn version 3.1
SEQ ID NO 1
LENGTH: 32
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:

OTHER INFORMATION: synthetic oligonucleotide
US-10-355-161A-1

Query Match 100.0%; Score 32; DB 17; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 9

US-10-831-778-429
Sequence 429, Application US/10831778
Publication No. US2004023474A1
GENERAL INFORMATION:
APPLICANT: Bratzler, Robert L.
APPLICANT: Petersen, Deanna M.
APPLICANT: Fournon, Yves
TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
FILE REFERENCE: C1037/7013 (HCL/MAT)
CURRENT APPLICATION NUMBER: US/10/831,778
CURRENT FILING DATE: 2004-04-23
PRIOR APPLICATION NUMBER: US 60/179,991
PRIOR FILING DATE: 2000-02-03
NUMBER OF SEQ ID NOS: 1093
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 429
LENGTH: 32
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Sequence
US-10-831-778-429

Query Match 100.0%; Score 32; DB 18; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.1e-07;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 10
US-09-931-583-35
Sequence 35, Application US/09931583
Publication No. US2003050263A1
GENERAL INFORMATION:
APPLICANT: Krieger, Arthur
APPLICANT: Kilman, Dennis
APPLICANT: Steinberg, Alfred
TITLE OF INVENTION: Methods and Products for Treating HIV Infection
FILE REFERENCE: C1039/7053 (HCL)
CURRENT APPLICATION NUMBER: US/09/931,583
CURRENT FILING DATE: 2001-08-16
PRIOR APPLICATION NUMBER: US 08/276,358
PRIOR FILING DATE: 1994-07-15
PRIOR APPLICATION NUMBER: US 09/415,142
PRIOR FILING DATE: 1999-10-09
NUMBER OF SEQ ID NOS: 75
SOFTWARE: PatentIn version 3.0
SEQ ID NO 35
LENGTH: 27
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
NAME/KEY: misc feature
OTHER INFORMATION: Synthetic Oligonucleotide
US-09-931-583-35

Query Match 81.2%; Score 26; DB 10; Length 27;
Best Local Similarity 100.0%; Pred. No. 0.00016;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCTTTGTGCTTTT 26
DB 1 TCGTCGTTTGTGCTTTGTGCTTTT 26

RESULT 11
US-09-888-326-131
; Sequence 131, Application US/09888326
; Publication No. US20030026801A1
; GENERAL INFORMATION:
; APPLICANT: Welner, George
; APPLICANT: Hartmann, Gunther
; TITLE OF INVENTION: Methods for Enhancing Antibody-Induced
; FILE REFERENCE: C1039/7052 (AMS)
; CURRENT APPLICATION NUMBER: US/09/888,326
; PRIOR FILING DATE: 2001-06-22
; NUMBER OF SEQ ID NOS: 848
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 131
; LENGTH: 29
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (0)...(0)
; OTHER INFORMATION: chimeric phosphorothioate/phosphodiester backbone
; NAME/KEY: misc_feature
; LOCATION: (1)...(1)
; OTHER INFORMATION: biotinylated at 5' end
US-09-888-326-131

Query Match 81.2%; Score 26; DB 10; Length 29;
Best Local Similarity 100.0%; Pred. No. 0.00016;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCTTTGTGCTTTT 26
DB 1 TCGTCGTTTGTGCTTTGTGCTTTT 26

RESULT 12
US-09-776-479-305
; Sequence 305, Application US/09776479
; Publication No. US20030087848A1
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; APPLICANT: Petersen, Deanna M.
; APPLICANT: Fournon, Yves
; TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
; FILE REFERENCE: C1037/7013 (HCL/MAT)
; CURRENT APPLICATION NUMBER: US/09/776,479
; PRIOR FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: US 60/179,991
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 305
; LENGTH: 29
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(3)

OTHER INFORMATION: Conjugated to biotin moiety.
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-09-776-479-305

Query Match 81.2%; Score 26; DB 10; Length 29;
Best Local Similarity 100.0%; Pred. No. 0.00016;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCTTTGTGCTTTT 26
DB 1 TCGTCGTTTGTGCTTTGTGCTTTT 26

RESULT 13
US-09-776-479-305
; Sequence 305, Application US/09776479
; Publication No. US20040067902A9
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; APPLICANT: Petersen, Deanna M.
; APPLICANT: Fournon, Yves
; TITLE OF INVENTION: Immunostimulatory Nucleic Acids for the
; FILE REFERENCE: C1037/7013 (HCL/MAT)
; CURRENT APPLICATION NUMBER: US/09/776,479
; PRIOR FILING DATE: 2001-02-02
; PRIOR APPLICATION NUMBER: US 60/179,991
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 305
; LENGTH: 29
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(3)
; OTHER INFORMATION: Conjugated to biotin moiety.
; FEATURE:
; OTHER INFORMATION: Synthetic Sequence
US-09-776-479-305

Query Match 81.2%; Score 26; DB 11; Length 29;
Best Local Similarity 100.0%; Pred. No. 0.00016;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCTTTGTGCTTTT 26
DB 1 TCGTCGTTTGTGCTTTGTGCTTTT 26

RESULT 14
US-10-112-653-295
; Sequence 295, Application US/10112653
; Publication No. US20030050268A1
; GENERAL INFORMATION:
; APPLICANT: Kriegl, Arthur M.
; APPLICANT: Berg, Daniel J.
; TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID FOR
; FILE REFERENCE: C01039/70060 (AMS)
; CURRENT APPLICATION NUMBER: US/10/112,653
; PRIOR FILING DATE: 2002-03-29
; PRIOR APPLICATION NUMBER: US 60/279,642
; NUMBER OF SEQ ID NOS: 1040
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 295
; LENGTH: 29
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:

OTHER INFORMATION: Synthetic Oligonucleotide
; NAME/KEY: misc.feature
; LOCATION: (1)...(1)
; OTHER INFORMATION: biotinylated
US-10-112-653-295

Query Match 81.2%; Score 26; DB 14; Length 29;
Best Local Similarity 100.0%; Pred. No. 0.00016;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTTT 26
|||
1 TCGTCGTTTGTGCGTTTGTGCGTTT 26

RESULT 15
US-10-017-995-305
; Sequence 305, Application US/10017995
; Publication No. US20030055014A1
; GENERAL INFORMATION:
; APPLICANT: Bratzler, Robert L.
; TITLE OF INVENTION: Inhibition of Angiogenesis by Nucleic Acids
; FILE REFERENCE: C1037/7025 (HCL/MAT)
; CURRENT APPLICATION NUMBER: US/10/017,995
; CURRENT FILING DATE: 2001-12-18
; PRIOR APPLICATION NUMBER: US 60/255,534
; PRIOR FILING DATE: 2000-12-14
; NUMBER OF SEQ ID NOS: 1093
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 305
; LENGTH: 29
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(3)
; OTHER INFORMATION: Conjugated to biotin moiety.
; OTHER INFORMATION: Synthetic Sequence
US-10-017-995-305

Query Match 81.2%; Score 26; DB 14; Length 29;
Best Local Similarity 100.0%; Pred. No. 0.00016;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTTT 26
|||
1 TCGTCGTTTGTGCGTTTGTGCGTTT 26

Search completed: February 14, 2005, 09:34:33
Job time : 303 secs

;; CURRENT FILING DATE: 2001-09-26
;; PRIOR APPLICATION NUMBER: US 09/082,649
;; PRIOR FILING DATE: 1998-05-20
;; PRIOR APPLICATION NUMBER: US 60/047,233
;; PRIOR FILING DATE: 1997-05-20
;; PRIOR APPLICATION NUMBER: US 60/047,209
;; NUMBER OF SEQ ID NOS: 84
;; SOFTWARE: FastSeq for Windows Version 3.0
;; SEQ ID NO 15
;; LENGTH: 52
;; TYPE: DNA
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: synthetic oligonucleotide
US-09-965-101-15

Query Match
Best Local Similarity 78.1%; Score 25; DB 4; Length 52;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTT 25
DB 4 TCGTCGTTTGTGCGTTTGTGCGTT 28

RESULT 3
US-09-030-701-6
; Sequence 6, Application US/09030701B
; Patent No. 6214806
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; APPLICANT: Schwartz, David A.
; TITLE OF INVENTION: USE OF NUCLEIC ACIDS CONTAINING
; TITLE OF INVENTION: UNMETHYLATED CGP DINUCLEOTIDE IN THE TREATMENT OF
; FILE REFERENCE: C1039/7011
; CURRENT APPLICATION NUMBER: US/09/030,701B
; PRIOR FILING DATE: 1998-02-25
; PRIOR APPLICATION NUMBER: 60/039,405
; NUMBER OF SEQ ID NOS: 65
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 6
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic oligonucleotide
US-09-030-701-6

Query Match
Best Local Similarity 75.0%; Score 24; DB 3; Length 24;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTT 24
DB 1 TCGTCGTTTGTGCGTTTGTGCGTT 24

RESULT 4
US-09-286-098-90
; Sequence 90, Application US/09286098
; Patent No. 6218371
; GENERAL INFORMATION:
; APPLICANT: Krieg, Arthur M.
; APPLICANT: Weiner, George
; TITLE OF INVENTION: Methods and Products for Stimulating the
; TITLE OF INVENTION: Immune System Using Immunotherapeutic Oligonucleotides and
; FILE REFERENCE: C1039/7026/HCT
; CURRENT APPLICATION NUMBER: US/09/286,098
; CURRENT FILING DATE: 1999-04-02

;; EARLIER APPLICATION NUMBER: US 60/080,729
;; EARLIER FILING DATE: 1998-04-03
;; NUMBER OF SEQ ID NOS: 105
;; SOFTWARE: FastSeq for Windows Version 3.0
;; SEQ ID NO 90
;; LENGTH: 24
;; TYPE: DNA
;; ORGANISM: Artificial Sequence
;; FEATURE:
;; OTHER INFORMATION: Synthetic Sequence
US-09-286-098-90

Query Match
Best Local Similarity 75.0%; Score 24; DB 3; Length 24;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTT 24
DB 1 TCGTCGTTTGTGCGTTTGTGCGTT 24

RESULT 5
US-08-960-774-46
; Sequence 46, Application US/08960774
; Patent No. 6239116
; GENERAL INFORMATION:
; APPLICANT: Krieg et al.,
; TITLE OF INVENTION: IMMUNOSTIMULATORY NUCLEIC ACID MOLECULES
; NUMBER OF SEQUENCES: 111
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 4225 Executive Square, Suite 1400
; CITY: La Jolla
; STATE: CA
; COUNTRY: USA
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: ASCII text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/960,774
; FILING DATE: 30-October-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 514
; FILING DATE: October 30, 1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Haile, Lisa A.
; REGISTRATION NUMBER: 38,347
; REFERENCE/DOCKET NUMBER: 08918/012001
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 619/678-5070
; TELEFAX: 619/678-5099
; INFORMATION FOR SEQ ID NO: 46:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 24 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
US-08-960-774-46

Query Match
Best Local Similarity 75.0%; Score 24; DB 3; Length 24;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTT 24
DB 1 TCGTCGTTTGTGCGTTTGTGCGTT 24

```
RESULT 6
US-09-082-649B-3
; Sequence 3, Application US/09082649B
; Patent No. 6339068
; GENERAL INFORMATION:
; APPLICANT: Davis, Heather L.
; APPLICANT: Kriegl, Arthur M.
; APPLICANT: Schorr, Joachim
; APPLICANT: Wu, Tong
; TITLE OF INVENTION: Vectors and Methods for Immunization or
; FILE REFERENCE: C1039/7009
; CURRENT APPLICATION NUMBER: US/09/082,649B
; PRIOR FILING DATE: 1998-05-20
; PRIOR APPLICATION NUMBER: US 60/047,233
; PRIOR FILING DATE: 1997-05-20
; PRIOR APPLICATION NUMBER: US 60/047,209
; PRIOR FILING DATE: 1997-05-20
; NUMBER OF SEQ ID NOS: 85
; SOFTWARE: FaastSeq for Windows Version 3.0
; SEQ ID NO 3
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic oligonucleotide
; NAME/KEY: misc_feature
; LOCATION: (0)..(0)
; OTHER INFORMATION: Has a phosphorothioate backbone.
US-09-082-649B-3

Query Match      75.0%; Score 24; DB 3; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00042;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 TCGTCGTTTGTGCTTTGTGCTT 24
Db      1 TCGTCGTTTGTGCTTTGTGCTT 24

RESULT 7
US-09-082-649B-66
; Sequence 66, Application US/09082649B
; Patent No. 6339068
; GENERAL INFORMATION:
; APPLICANT: Davis, Heather L.
; APPLICANT: Kriegl, Arthur M.
; APPLICANT: Schorr, Joachim
; APPLICANT: Wu, Tong
; TITLE OF INVENTION: Vectors and Methods for Immunization or
; FILE REFERENCE: C1039/7009
; CURRENT APPLICATION NUMBER: US/09/082,649B
; PRIOR FILING DATE: 1998-05-20
; PRIOR APPLICATION NUMBER: US 60/047,233
; PRIOR FILING DATE: 1997-05-20
; PRIOR APPLICATION NUMBER: US 60/047,209
; PRIOR FILING DATE: 1997-05-20
; NUMBER OF SEQ ID NOS: 85
; SOFTWARE: FaastSeq for Windows Version 3.0
; SEQ ID NO 66
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: synthetic oligonucleotide
; NAME/KEY: misc_feature
; LOCATION: (0)..(0)
; OTHER INFORMATION: Backbone is a phosphorothioate--phosphodiester
; OTHER INFORMATION: chimera.
US-09-082-649B-66
```

```
Query Match      75.0%; Score 24; DB 3; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00042;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 TCGTCGTTTGTGCTTTGTGCTT 24
Db      1 TCGTCGTTTGTGCTTTGTGCTT 24

RESULT 8
US-09-325-193A-77
; Sequence 77, Application US/09325193A
; Patent No. 6406705
; GENERAL INFORMATION:
; APPLICANT: Davis, Heather L.
; APPLICANT: Schorr, Joachim
; APPLICANT: Kriegl, Arthur M.
; TITLE OF INVENTION: Use of Nucleic Acids Containing
; FILE REFERENCE: C1039/7025/HCL
; CURRENT APPLICATION NUMBER: US/09/325,193A
; PRIOR FILING DATE: 1999-06-03
; PRIOR APPLICATION NUMBER: US 09/154,614
; PRIOR FILING DATE: 1998-09-16
; PRIOR APPLICATION NUMBER: PCT/US98/04703
; PRIOR FILING DATE: 1998-03-10
; PRIOR APPLICATION NUMBER: US 60/040,376
; PRIOR FILING DATE: 1997-03-10
; NUMBER OF SEQ ID NOS: 98
; SOFTWARE: FaastSeq for Windows Version 3.0
; SEQ ID NO 77
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Synthetic Oligonucleotide
US-09-325-193A-77

Query Match      75.0%; Score 24; DB 3; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00042;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      1 TCGTCGTTTGTGCTTTGTGCTT 24
Db      1 TCGTCGTTTGTGCTTTGTGCTT 24

RESULT 9
US-09-191-170-84
; Sequence 84, Application US/09191170
; Patent No. 6429199
; GENERAL INFORMATION:
; APPLICANT: Kriegl, Arthur M.
; APPLICANT: Hartmann, Gunther
; TITLE OF INVENTION: Immunostimulatory Nucleic Acid Molecules
; FILE REFERENCE: C1039/7017
; CURRENT APPLICATION NUMBER: US/09/191,170
; PRIOR FILING DATE: 1998-11-13
; PRIOR APPLICATION NUMBER: US 08/960,774
; PRIOR FILING DATE: 1997-10-30
; PRIOR APPLICATION NUMBER: US 08/738,652
; PRIOR FILING DATE: 1996-10-30
; PRIOR APPLICATION NUMBER: US 08/386,063
; PRIOR FILING DATE: 1995-02-07
; PRIOR APPLICATION NUMBER: US 08/276,358
; PRIOR FILING DATE: 1994-07-15
; NUMBER OF SEQ ID NOS: 99
; SOFTWARE: FaastSeq for Windows Version 3.0
; SEQ ID NO 84
; LENGTH: 24
; TYPE: DNA
; ORGANISM: Artificial Sequence
```

FEATURE:
OTHER INFORMATION: synthetic oligonucleotide
US-09-191-170-84

Query Match
Best Local Similarity 75.0%; Score 24; DB 3; Length 24;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCGTCGTTTGTGCTTTGTCGTT 24
Db 1 TCGTCGTTTGTGCTTTGTCGTT 24

RESULT 10
US-09-191-170-95

Sequence 95, Application US/09191170
Patent No. 6429199
GENERAL INFORMATION:
APPLICANT: Kriegl, Arthur M.
APPLICANT: Hartmann, Gunther
TITLE OF INVENTION: Immunostimulatory Nucleic Acid Molecules
FILE REFERENCE: C1039/7017
CURRENT APPLICATION NUMBER: US/09/191,170
EARLIER FILING DATE: 1998-11-13
EARLIER APPLICATION NUMBER: US 08/960,774
EARLIER FILING DATE: 1997-10-30
EARLIER APPLICATION NUMBER: US 08/738,652
EARLIER FILING DATE: 1996-10-30
EARLIER APPLICATION NUMBER: US 08/386,063
EARLIER FILING DATE: 1995-02-07
EARLIER APPLICATION NUMBER: US 08/276,358
EARLIER FILING DATE: 1994-07-15
NUMBER OF SEQ ID NOS: 99
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 95

LENGTH: 24
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: synthetic oligonucleotide
FEATURE:
NAME/KEY: modified base
LOCATION: (2)...(2)
OTHER INFORMATION: m5c
FEATURE:
NAME/KEY: modified base
LOCATION: (5)...(5)
OTHER INFORMATION: m5c
FEATURE:
NAME/KEY: modified base
LOCATION: (13)...(13)
OTHER INFORMATION: m5c
FEATURE:
NAME/KEY: modified base
LOCATION: (21)...(21)
OTHER INFORMATION: m5c
US-09-191-170-95

Query Match
Best Local Similarity 75.0%; Score 24; DB 3; Length 24;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCGTCGTTTGTGCTTTGTCGTT 24
Db 1 TCGTCGTTTGTGCTTTGTCGTT 24

RESULT 11
US-09-690-921-4

Sequence 4, Application US/09690921
Patent No. 6544518
GENERAL INFORMATION:

APPLICANT: Friede, Martin
APPLICANT: Gerard, Catherine
APPLICANT: Hermand, Philippe
TITLE OF INVENTION: Vaccines
FILE REFERENCE: B45181-1
CURRENT APPLICATION NUMBER: US/09/690,921
CURRENT FILING DATE: 2000-10-18
PRIOR APPLICATION NUMBER: PCT/EP00/02920
PRIOR FILING DATE: 2000-04-04
PRIOR APPLICATION NUMBER: 09/301,829
PRIOR FILING DATE: 1999-04-29
PRIOR APPLICATION NUMBER: 990885.8
PRIOR FILING DATE: 1999-04-19
NUMBER OF SEQ ID NOS: 5
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 4

LENGTH: 24
TYPE: DNA
ORGANISM: Human
US-09-690-921-4

Query Match
Best Local Similarity 75.0%; Score 24; DB 4; Length 24;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCGTCGTTTGTGCTTTGTCGTT 24
Db 1 TCGTCGTTTGTGCTTTGTCGTT 24

RESULT 12
US-09-337-619-46

Sequence 46, Application US/09337619
Patent No. 6653292
GENERAL INFORMATION:
APPLICANT: Kriegl, Arthur M.
TITLE OF INVENTION: Methods of Treating Cancer Using
FILE REFERENCE: C1039/7021/HCL
CURRENT APPLICATION NUMBER: US/09/337,619
CURRENT FILING DATE: 1999-06-21
EARLIER FILING DATE: 1997-10-30
EARLIER APPLICATION NUMBER: US 08/960,774
EARLIER FILING DATE: 1997-10-30
EARLIER APPLICATION NUMBER: US 08/738,652
EARLIER FILING DATE: 1996-10-30
EARLIER APPLICATION NUMBER: US 08/386,063
EARLIER FILING DATE: 1995-02-07
EARLIER APPLICATION NUMBER: US 08/276,358
EARLIER FILING DATE: 1994-07-15
NUMBER OF SEQ ID NOS: 123
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 46

LENGTH: 24
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Synthetic Oligonucleotide
US-09-337-619-46

Query Match
Best Local Similarity 75.0%; Score 24; DB 4; Length 24;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCGTCGTTTGTGCTTTGTCGTT 24
Db 1 TCGTCGTTTGTGCTTTGTCGTT 24

RESULT 13
US-09-965-101-3

Sequence 3, Application US/09965101
Patent No. 6821957
GENERAL INFORMATION:

```
/ APPLICANT: Davis, Heather L.
/ APPLICANT: Kriegl, Arthur M.
/ APPLICANT: Schorr, Joachim
/ APPLICANT: Wu, Tong
/ TITLE OF INVENTION: Vectors and Methods for Immunization or
/ FILE REFERENCE: C1039/7057 (HCL/MAT)
/ CURRENT APPLICATION NUMBER: US/09/965,101
/ CURRENT FILING DATE: 2001-09-26
/ PRIOR APPLICATION NUMBER: US 09/082,649
/ PRIOR FILING DATE: 1998-05-20
/ PRIOR APPLICATION NUMBER: US 60/047,233
/ PRIOR FILING DATE: 1997-05-20
/ PRIOR APPLICATION NUMBER: US 60/047,209
/ PRIOR FILING DATE: 1997-05-20
/ NUMBER OF SEQ ID NOS: 84
/ SOFTWARE: FastSeq for Windows Version 3.0
/ SEQ ID NO 3
/ LENGTH: 24
/ TYPE: DNA
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: synthetic oligonucleotide
/ NAME/KEY: misc_feature
/ LOCATION: (0)..(0)
/ OTHER INFORMATION: Has a phosphorothioate backbone.
US-09-965-101-3
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Query Match          75.0%; Score 24; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00042;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      1 TCGTCGTTTGTGCTTTGTCGTT 24
Db      1 TCGTCGTTTGTGCTTTGTCGTT 24
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RESULT 14
US-09-965-101-66
/ Sequence 66, Application US/09965101
/ Patent No. 6821957
/ GENERAL INFORMATION:
/ APPLICANT: Davis, Heather L.
/ APPLICANT: Kriegl, Arthur M.
/ APPLICANT: Schorr, Joachim
/ APPLICANT: Wu, Tong
/ TITLE OF INVENTION: Vectors and Methods for Immunization or
/ FILE REFERENCE: C1039/7057 (HCL/MAT)
/ CURRENT APPLICATION NUMBER: US/09/965,101
/ CURRENT FILING DATE: 2001-09-26
/ PRIOR APPLICATION NUMBER: US 09/082,649
/ PRIOR FILING DATE: 1998-05-20
/ PRIOR APPLICATION NUMBER: US 60/047,233
/ PRIOR FILING DATE: 1997-05-20
/ PRIOR APPLICATION NUMBER: US 60/047,209
/ PRIOR FILING DATE: 1997-05-20
/ NUMBER OF SEQ ID NOS: 84
/ SOFTWARE: FastSeq for Windows Version 3.0
/ SEQ ID NO 66
/ LENGTH: 24
/ TYPE: DNA
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: synthetic oligonucleotide
/ NAME/KEY: misc_feature
/ LOCATION: (0)..(0)
/ OTHER INFORMATION: Backbone is a phosphorothioate--phosphodiester
/ OTHER INFORMATION: chimera.
US-09-965-101-66
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Query Match          75.0%; Score 24; DB 4; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00042;
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Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      1 TCGTCGTTTGTGCTTTGTCGTT 24
Db      1 TCGTCGTTTGTGCTTTGTCGTT 24
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RESULT 15
US-09-337-619-123
/ Sequence 123, Application US/09337619
/ Patent No. 6653292
/ GENERAL INFORMATION:
/ APPLICANT: Kriegl, Arthur M.
/ TITLE OF INVENTION: Methods of Treating Cancer Using
/ FILE REFERENCE: C1039/7021/HCL
/ CURRENT APPLICATION NUMBER: US/09/337,619
/ CURRENT FILING DATE: 1999-06-21
/ EARLIER APPLICATION NUMBER: US 08/960,774
/ EARLIER FILING DATE: 1997-10-30
/ EARLIER APPLICATION NUMBER: US 08/738,652
/ EARLIER FILING DATE: 1996-10-30
/ EARLIER APPLICATION NUMBER: US 08/386,063
/ EARLIER FILING DATE: 1995-02-07
/ EARLIER APPLICATION NUMBER: US 08/276,358
/ EARLIER FILING DATE: 1994-07-15
/ NUMBER OF SEQ ID NOS: 123
/ SOFTWARE: FastSeq for Windows Version 3.0
/ SEQ ID NO 123
/ LENGTH: 23
/ TYPE: DNA
/ ORGANISM: Artificial Sequence
/ FEATURE:
/ OTHER INFORMATION: Synthetic Oligonucleotide
US-09-337-619-123
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Query Match          71.9%; Score 23; DB 4; Length 23;
Best Local Similarity 100.0%; Pred. No. 0.0014;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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QY      1 TCGTCGTTTGTGCTTTGTCGTT 23
Db      1 TCGTCGTTTGTGCTTTGTCGTT 23
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Search completed: February 14, 2005, 07:23:18
Job time : 103 secs
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REFERENCE 1
AUTHORS Weiner, G. and Hartmann, G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
JOURNAL Cancer
PATENT: WO 0197843-A 734 29-DEC-2001
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)
FEATURES
SOURCE
1. .32
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic oligonucleotide-phosphorothioate backbone"

ORIGIN

Query Match 100.0%; Score 32; DB 6; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.3e-08;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 3

LOCUS AX547290 32 bp DNA linear PAT 01-MAR-2003
DEFINITION Sequence 429 from Patent WO02053141.
ACCESSION AX547290
VERSION AX547290.1 GI:25812434
KEYWORDS
SOURCE
ORGANISM
synthetic construct
synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Bratzler, R.L.
TITLE Inhibition of angiogenesis by nucleic acids
JOURNAL Patent: WO 02053141-A 429 11-JUL-2002;
Coley Pharmaceutical Group, Inc. (US)
FEATURES
SOURCE
1. .32
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic Sequence"

ORIGIN

Query Match 100.0%; Score 32; DB 6; Length 32;
Best Local Similarity 100.0%; Pred. No. 1.3e-08;
Matches 32; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32
1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 32

RESULT 4

LOCUS AX104113 29 bp DNA linear PAT 30-APR-2001
DEFINITION Sequence 305 from Patent WO0122972.
ACCESSION AX104113
VERSION AX104113.1 GI:13920310
KEYWORDS
SOURCE
ORGANISM
synthetic construct
synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Kriegl, A.M., Schetter, C. and Vollmer, J.C.
TITLE Immuno-stimulatory nucleic acids
JOURNAL Patent: WO 0122972-A 305 05-APR-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US); Coley Pharmaceutical
GmbH (DE)
FEATURES
SOURCE
Location/Qualifiers

source

1. .29
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Biotin moiety attached at 5' end of sequence."

QY 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 26
1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 26

Db 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 26

RESULT 5

LOCUS AX355103 29 bp DNA linear PAT 06-FEB-2002
DEFINITION Sequence 131 from Patent WO0197843.
ACCESSION AX355103
VERSION AX355103.1 GI:18619770
KEYWORDS
SOURCE
ORGANISM
synthetic construct
synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Weiner, G. and Hartmann, G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
JOURNAL Cancer
PATENT: WO 0197843-A 131 27-DEC-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)
FEATURES
SOURCE
1. .29
/organism="synthetic construct"
/mol_type="unassigned DNA"
/db_xref="taxon:32630"
/note="Synthetic oligonucleotide-chimeric phosphorothioate/phosphodiester backbone with phosphodiester on 5' end"

ORIGIN

Query Match 81.2%; Score 26; DB 6; Length 29;
Best Local Similarity 100.0%; Pred. No. 4.4e-05;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 26
1 TCGTCGTTTGTGCGTTTGTGCGTTTGTGCGTT 26

RESULT 6

LOCUS AX547166 29 bp DNA linear PAT 01-MAR-2003
DEFINITION Sequence 305 from Patent WO02053141.
ACCESSION AX547166
VERSION AX547166.1 GI:25812310
KEYWORDS
SOURCE
ORGANISM
synthetic construct
synthetic construct
other sequences; artificial sequences.

REFERENCE 1
AUTHORS Bratzler, R.L.
TITLE Inhibition of angiogenesis by nucleic acids
JOURNAL Patent: WO 02053141-A 305 11-JUL-2002;
Coley Pharmaceutical Group, Inc. (US)
FEATURES
SOURCE
1. .29
/organism="synthetic construct"
/mol_type="unassigned DNA"

/db_xref="taxon:32630"
misc_feature 1..3
/note="Conjugated to biotin moiety.
Synthetic Sequence"
ORIGIN
Query Match 81.2%; Score 26; DB 6; Length 29;
Best Local Similarity 100.0%; Pred. No. 4.4e-05;
Matches 26; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCGTCGTTTGTGCGTTTGTGCGTTT 26
1 TCGTCGTTTGTGCGTTTGTGCGTTT 26
Db 1 TCGTCGTTTGTGCGTTTGTGCGTTT 26
RESULT 7
ARI82843 52 bp DNA linear PAT 20-APR-2002
LOCUS Sequence 15 from patent US 6339068.
DEFINITION ARI82843
ACCESSION ARI82843 GI:20226050
VERSION ARI82843.1 GI:20226050
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 52)
AUTHORS Kriegl,A.M., Davis,H.L., Wu,T. and Schorr,J.
TITL E Vectors and methods for immunization or therapeutic protocols
JOURNAL Patent: US 6339068-A 15 15-JAN-2002;
FEATURES location/Qualifiers
1..52
/organism="unknown"
/mol_type="unassigned DNA"
ORIGIN
Query Match 78.1%; Score 25; DB 6; Length 52;
Best Local Similarity 100.0%; Pred. No. 0.00018;
Matches 25; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCGTCGTTTGTGCGTTTGTGCGTTT 25
4 TCGTCGTTTGTGCGTTTGTGCGTTT 28
Db 4 TCGTCGTTTGTGCGTTTGTGCGTTT 28
RESULT 8
ARI46378 24 bp DNA linear PAT 08-AUG-2001
LOCUS Sequence 90 from patent US 6218371.
DEFINITION ARI46378
ACCESSION ARI46378
VERSION ARI46378.1 GI:15109567
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 24)
AUTHORS Kriegl,A.M. and Weiner,G.
TITL E Methods and products for stimulating the immune system using
JOURNAL immunotherapeutic oligonucleotides and cytokines
PATent: US 6218371-A 90 17-APR-2001;
FEATURES location/Qualifiers
1..24
/organism="unknown"
/mol_type="unassigned DNA"
ORIGIN
Query Match 75.0%; Score 24; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00066;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCGTCGTTTGTGCGTTTGTGCGTT 24
1 TCGTCGTTTGTGCGTTTGTGCGTT 24
Db 1 TCGTCGTTTGTGCGTTTGTGCGTT 24

RESULT 9
ARI54717 24 bp DNA linear PAT 08-AUG-2001
LOCUS Sequence 46 from patent US 6239116.
DEFINITION ARI54717
ACCESSION ARI54717
VERSION ARI54717.1 GI:15122770
KEYWORDS
SOURCE Unknown.
ORGANISM Unknown.
REFERENCE 1 (bases 1 to 24)
AUTHORS Kriegl,A.M. and Kline,J.N.
TITL E Immunostimulatory nucleic acid molecules
JOURNAL Patent: US 6239116-A 46 29-MAY-2001;
FEATURES location/Qualifiers
1..24
/organism="unknown"
/mol_type="unassigned DNA"
ORIGIN
Query Match 75.0%; Score 24; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00066;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCGTCGTTTGTGCGTTTGTGCGTT 24
1 TCGTCGTTTGTGCGTTTGTGCGTT 24
Db 1 TCGTCGTTTGTGCGTTTGTGCGTT 24
RESULT 10
BD205600 24 bp DNA linear PAT 17-JUL-2003
LOCUS Method of controlling hematopoiesis by using CpG oligonucleotide.
DEFINITION BD205600
ACCESSION BD205600.1 GI:33015370
VERSION BD205600.1 GI:33015370
KEYWORDS JP 2002514397-A/90.
SOURCE JP 2002514397-A/90.
ORGANISM
REFERENCE 1 (bases 1 to 24)
AUTHORS Wagner,H. and Lipford,G.
TITL E Method of controlling hematopoiesis by using CpG oligonucleotide
JOURNAL Patent: JP 2002514397-A 90 21-MAY-2002;
COMMENT CORY PHARMACEUTICALS GMBH,CORY PHARMACEUTICALS GROUP INC
OS Artificial Sequence
PN JP 2002514397-A/90
PD 21-MAY-2002
PF 14-MAY-1999 JP 2000547969
PR 14-MAY-1998 US 60/085516,02-FEB-1999 US 09/241653 PT
HERMANN WAGNER,GRAYSON LIPFORD
PC C12N15/09,A61K31/70,A61K39/39,C07H21/04//A61K45/00,C12N15/00
CC Synthetic Sequence
FH Key location/Qualifiers
FT source 1..24
/organism="Artificial Sequence".
FEATURES location/Qualifiers
1..24
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
ORIGIN
Query Match 75.0%; Score 24; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00066;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCGTCGTTTGTGCGTTTGTGCGTT 24
1 TCGTCGTTTGTGCGTTTGTGCGTT 24
Db 1 TCGTCGTTTGTGCGTTTGTGCGTT 24
RESULT 11

BD261142 24 bp DNA linear PAT 17-JUL-2003
LOCUS
DEFINITION Method and products for stimulating the immune system using
immunotherapeutic oligonucleotides and cytokines.
ACCESSION BD261142
VERSION BD261142.1 GI:33070912
KEYWORDS JP 2002510644-A/90.
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 24)
AUTHORS Kriegl,A.M. and Weiner,G.
TITLE Methods and products for stimulating the immune system using
immunotherapeutic oligonucleotides and cytokines
JOURNAL Patent: JP 2002510644-A 90 09-APR-2002,
UNIVERSITY OF IOWA RESEARCH FOUNDATION
COMMENT OS Artificial Sequence
PN JP 2002510644-A/90
PD 09-APR-2002
PF 02-APR-1999 JP 2000542030
PR 03-APR-1998 US 60/080729
PI ARTHUR M KRIEGL,GEORGE WEINER
PC A61K38/00,A61K31/7088,A61K39/00,A61P15/00,A61P35/00,A61P37/04,
CC A61K37/02
CC Synthetic Sequence
FT Key Location/Qualifiers
FT source 1..24
/organism='Artificial Sequence'.
FEATURES
source 1..24
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"
ORIGIN
Query Match 75.0%; Score 24; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00066;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCGTCGTTTGTGCTTTGTCGTT 24
1 TCGTCGTTTGTGCTTTGTCGTT 24
Db 1 TCGTCGTTTGTGCTTTGTCGTT 24
RESULT 12 24 bp DNA linear PAT 17-JUL-2003
BD261298
LOCUS
DEFINITION Methods and products for inducing mucosal immunity.
ACCESSION BD261298
VERSION BD261298.1 GI:33071068
KEYWORDS JP 2002516294-A/77.
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 24)
AUTHORS McCluskie,M.J. and Davis,H.L.
TITLE Methods and products for inducing mucosal immunity
JOURNAL Patent: JP 2002516294-A 77 04-JUN-2002,
LOEB HEALTH RESEARCH INSTITUTE AT THE OTTAWA HOSPITAL, CORY
PHARMACEUTICALS GROUP INC
COMMENT OS Artificial Sequence
PN JP 2002516294-A/77
PD 04-JUN-2002
PF 21-MAY-1999 JP 2000550515
PR 22-MAY-1998 US 60/086393
PI MICHAEL J MCCCLUSKIE, HEATHER L DAVIS
PC A61K39/00,A61K9/10,A61K9/16,A61K9/50,A61K9/51,A61K31/70,A61K39/ PC
39, A61P31/00,A61P35/00,A61P37/00
CC immunostimulatory synthetic oligonucleotide
FH Key Location/Qualifiers
FT source 1..24

BD261563 24 bp DNA linear PAT 17-JUL-2003
LOCUS
DEFINITION Vaccine.
ACCESSION BD261563
VERSION BD261563.1 GI:33071331
KEYWORDS JP 2002542203-A/4.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 24)
AUTHORS Friede,M., Garcon,N. and Hermand,P.
TITLE Vaccine
JOURNAL Patent: JP 2002542203-A 4 10-DEC-2002;
SMITHKLINE BEECHAM BIOLOGICALS SA
COMMENT OS Homo sapiens (human)
PN JP 2002542203-A/4
PD 10-DEC-2002
PF 04-APR-2000 JP 2000611936
PR 19-APR-1999 GB 9908885.8,29-APR-1999 US 09/301829 PI
MARTIN FRIEDE,NATHALIE GARCON,PHILIPPE HERMAND PC
A61K39/39,A61K31/7088,A61K39/00,A61K39/00,A61K39/02, PC
A61K39/095,
PC A61K39/10,A61K39/102,A61K39/112,A61K39/118,A61K39/12,A61K39/
PC 145,A61K39/21,
PC A61K39/245,A61K39/25,A61K39/29,A61P9/10,A61P25/28,A61P31/04,
PC A61P31/12,
PC A61P33/00,A61P33/02,A61P35/00,A61P37/04,A61P37/08,A61P43/00,
PC C12N15/09,
CC C12N15/00
CC Vaccine
FH Key Location/Qualifiers
FT source 1..24
/organism='Homo sapiens (human)'.
FEATURES
source 1..24
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
ORIGIN
Query Match 75.0%; Score 24; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00066;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCGTCGTTTGTGCTTTGTCGTT 24
1 TCGTCGTTTGTGCTTTGTCGTT 24
Db 1 TCGTCGTTTGTGCTTTGTCGTT 24
RESULT 14 24 bp DNA linear PAT 17-JUL-2003
BD267904
LOCUS
DEFINITION Methods for the prevention and treatment of parasitic infections
and related diseases using CPG oligonucleotides.

ACCESSION BD267904
VERSION BD267904.1 GI:33077672
KEYWORDS JP 2002513763-A/77.
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 24)
AUTHORS Gramaneti R.A., Krieg,A.M., Davis,H.L. and Hoffman,S.L.
TITLE Methods for the prevention and treatment of parasitic infections
and related diseases using CPG oligonucleotides
JOURNAL Patent: JP 2002513763-A 77 14-MAY-2002;
UNIVERSITY OF IOWA RESEARCH FOUNDATION, OTTAWA CIVIC LOEB RESEARCH
INSTITUTE, UNITED STATES OF AMERICA AS REPRESENTED BY THE SECRETARY
OF THE NAVY
COMMENT OS Artificial Sequence
PN JP 2002513763-A/77
PD 14-MAY-2002
PR 06-MAY-1999 JP 2000546780
PI 06-MAY-1998 US 60/084512
PI ROBERT A GRAMANETI,ARTHUR M KRIEG,HEATHER L DAVIS,STEPHEN L
PI HOFFMAN
PC A61K31/711,A61K9/127,A61K38/00,A61K38/22,A61K45/00,A61P31/00,
PC A61P33/00//
PC C12N15/09,A61K37/02,A61K37/24,C12N15/00
CC Synthetic Sequence
FH Key Location/Qualifiers
FT source 1..24
Location/Qualifiers
1..24
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"

ORIGIN
Query Match 75.0%; Score 24; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00066;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TCGTCGTTTGTGCGTTTGTGCTT 24
1 TCGTCGTTTGTGCGTTTGTGCTT 24

Db
1 TCGTCGTTTGTGCGTTTGTGCTT 24

RESULT 15
BD270804 24 bp DNA linear PAT 17-JUL-2003
LOCUS BD270804
DEFINITION Stereoisomer of Cpg oligonucleotide and method relating thereto.
ACCESSION BD270804
VERSION BD270804.1 GI:33080572
KEYWORDS JP 2002521489-A/77.
SOURCE synthetic construct
ORGANISM synthetic construct
other sequences; artificial sequences.
REFERENCE 1 (bases 1 to 24)
AUTHORS Krieg,A.M.
TITLE Stereoisomer of Cpg oligonucleotide and method relating thereto
JOURNAL Patent: JP 2002521489-A 77 16-JUL-2002;
UNIVERSITY OF IOWA RESEARCH FOUNDATION
COMMENT OS Artificial Sequence
PN JP 2002521489-A/77
PD 16-JUL-2002
PR 27-JUL-1999 JP 2000562385
PR 27-JUL-1998 US 60/094370
PI ARTHUR M KRIEG
PC A61K31/711,A61P11/06,A61P17/00,A61P27/02,A61P29/00,A61P31/00,
PC A61P33/00,
PC A61P35/00,A61P37/04,A61P37/06,A61P37/08
CC Synthetic
FH Key Location/Qualifiers
FT source 1..24
Location/Qualifiers
1..24
/organism="Artificial Sequence".

FEATURES
Location/Qualifiers

SOURCE

1..24
/organism="synthetic construct"
/mol_type="genomic DNA"
/db_xref="taxon:32630"

ORIGIN

Query Match 75.0%; Score 24; DB 6; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.00066;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGTGCGTTTGTGCTT 24
1 TCGTCGTTTGTGCGTTTGTGCTT 24

Db

Search completed: February 14, 2005, 07:21:31
Job time : 1727 secs

Submitted (31-MAR-2004) Weishaar B., Max-Planck-Institut fuer Zuechtungsforshung, Carl-Neuberg-Strasse 11, 50829, Germany. This sequence has been recovered from the right border of the T-DNA. Details on the protocols used for generation of the sequence are described in References 1-3. Re-examination of the source from which this sequence has been produced indicates that the sequence is of low reliability. Therefore, no information on a potential insertion site is deduced. The sequences are generated at the MPI

for Plant Breeding Research in the context of the GABI-Kat project.
GABI-Kat is part of the German plant Genomics program designated
'GABI'. Information on line availability can be found at:
<http://www.mpiz-koeln.mpg.de/GABI-Kat/>.
Location/Qualifiers
1. .38

FEATURES
source
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Colombia 0"
/db_xref="taxon:3702"
/clone="GK-106B12-012499"
/clone_1lb="Arabidopsis thaliana T-DNA insertion lines"
/ecotype="Col-0"
/note="PCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector PAC161 (Genbank accession number: AJ537514). The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. T-DNA derived sequences were removed."

ORIGIN
Query Match
Best Local Similarity 43.8%; Score 14; DB 9; Length 38;
Matches 14; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Db 7 TTTTGTGCTTTTGT 20
29 TTTTGTGCTTTTGT 16

RESULT 2
B0613522/c
LOCUS
DEFINITION
r008f05.v1 Meloidogyne incognita egg SL1 TOPO v1 Meloidogyne
ACCESSION
B0613522
VERSION
B0613522.1 GI:21603191
KEYWORDS
EST.
SOURCE
ORGANISM
Meloidogyne incognita (southern root-knot nematode)
Eukaryota; Metazoa; Nematoda; Chromadorea; Tylenchida; Tylenchida;
Tylenchoidea; Heterodidae; Meloidogyninae; Meloidogyne.
1 (bases 1 to 50)
McCartter,J., Clifton,S., Chapelli,B., Page,D., Martin,J.,
Wyle,T., Dante,M., Marra,M., Hillier,L., Kucaba,T., Theising,B.,
Bowers,Y., Gibbons,M., Ritter,F., Bennett,D., Franklin,C.,
Tasgareishvili,R., Ronko,I., Kennedy,S., Maguire,L., Beck,C.,
Harvey,N., Schurk,R., Kohn,S., Shin,T., Jackson,Y., Cardenas,M.,
McCam,R., Waterston,R. and Wilson,R.
The Washington Univ. Nematode EST Project, 1999
Unpublished (1999)
Contact: McCartter JP
The Washington Univ. Nematode EST Project, 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.wustl.edu
The library was constructed by Claire Murphy and Dr. James McCartter
at Washington University, St. Louis. Meloidogyne incognita eggs
were provided by Andrew Kloek of Divergence Inc., St. Louis, MO.
The vector to vector length read
Seq primer: -40bp from Gibco.

FEATURES
source
Location/Qualifiers
1. .50
/organism="Meloidogyne incognita"
/mol_type="mRNA"
/db_xref="taxon:6306"
/dev_stage="egg"
/lab_host="DH10B (Invitrogen)"

/clone_1lb="Meloidogyne incognita egg SL1 TOPO v1"
/note="Vector: PCR1-TOPO (Invitrogen); Site 1: EcoRI,
Site 2: EcoRI; The library was constructed by Claire
Murphy and Dr. James McCartter at Washington University,
St. Louis. Oligo(dT)-SL1 PCR based library. cDNA PCR
products of size >400 nucleotides containing SL1 on the 5'
end and oligo(dT) on the 3' end were non-directionally
cloned into PCR1-TOPO (Invitrogen) following the TOPO TA
cloning protocol. Meloidogyne incognita eggs were provided
by Andrew Kloek of Divergence Inc., St. Louis, MO."

ORIGIN
Query Match
Best Local Similarity 40.6%; Score 13; DB 5; Length 50;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Db 6 GTTTGTGCTTTT 18
15 GTTTGTGCTTTT 3

RESULT 3
A0600996/c
LOCUS
DEFINITION
Arabidopsis thaliana T-DNA flanking sequence, right border, clone
518D07, genomic survey sequence.
ACCESSION
A0600996
VERSION
A0600996.1 GI:37950624
KEYWORDS
GSS; right border; T-DNA flanking sequence.
SOURCE
Arabidopsis thaliana (thale cress)
ORGANISM
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.

REFERENCE
AUTHORS
1 Brunaud,V., Balzerque,S., Dubreucq,B., Aubourg,S., Samson,F.,
Chauvin,S., Bechtold,N., Crnaud,C., Derose,R., Pelleter,G.,
Lepiniec,L., Caboche,M. and Lecharny,A.
T-DNA integration into the Arabidopsis genome depends on sequences
of pre-insertion sites
EMBO Rep. 3 (12), 1152-1157 (2002)
22363535
12445655
2 (bases 1 to 23)
Balzerque,S.
Direct Submission
Submitted (23-OCT-2003) Balzerque S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
PCR was performed on DNA from transformants of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
<http://dbsgap.versailles.inra.fr/publications/>. This sequence has
been generated in the framework of the French plant genomics
program 'Genoplante' (<http://www.genoplante.com> and
<http://genoplante.info.inbioigen.fr>).
Location/Qualifiers
1. .23

FEATURES
source
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/cultivar="Wassilewskija"
/db_xref="taxon:3702"
/clone="518D07"
/clone_1lb="Arabidopsis thaliana T-DNA insertion lines"
/note="T-DNA flanking sequence
right border"

ORIGIN
Query Match
37.5%; Score 12; DB 9; Length 23;

Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 TTTTGTGCTTTT 18
|||||
15 TTTTGTGCTTTT 4

Db 15 TTTTGTGCTTTT 4

RESULT 4
AJ595925 34 bp DNA linear GSS 15-JAN-2004
LOCUS Arabidopsis thaliana T-DNA flanking sequence, left border, clone
DEFINITION 425H07, genomic survey sequence.
ACCESSION AJ595925
VERSION AJ595925
KEYWORDS GSS; left border; T-DNA flanking sequence.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1
Brunaud, V., Balzerque, S., Dubreucq, B., Aubourg, S., Samson, P.,
Chauvin, S., Bechtold, N., Cruaud, C., DeRose, R., Pelletier, G.,
Lepoint, L., Caboche, M. and Lecharny, A.
T-DNA integration into the Arabidopsis genome depends on sequences
of pre-insertion sites
EMBO Rep. 3 (12), 1152-1157 (2002)

JOURNAL MEDLINE
22363535
PUBMED 12446565

REFERENCE 2 (bases 1 to 34)
Balzerque, S.
Direct Submission
Submitted (23-OCT-2003) Balzerque S., UMRGV, INRA/CNRS, 2 rue
Gaston Cremieux, 91057 Evry cedex, FRANCE
PCR was performed on DNA from transfections of Arabidopsis thaliana
plants from INRA (Versailles). The DNA fragment(s) resulting from
the PCR were directly sequenced from the left or the right border
to determine the genomic sequence flanking the insertion. T-DNA
derived sequences were removed. Information to order the
corresponding mutant line and a link to a database providing a
graphical display of the insertion site are available at
http://dbsg.versailles.inra.fr/publiclines/. This sequence has
been generated in the framework of the French plant genomics
program "Genoplante" (http://www.genoplante.com and
http://genoplante-info.inbio.gen.fr).

FEATURES
source
1..34
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/cultiivar="Wassiljewskij4"
/db_xref="taxon:3702"
/clone_1ib="425H07"
/clone_1ib="Arabidopsis thaliana T-DNA insertion lines"
1..34
/note="T-DNA flanking sequence
left border"

ORIGIN
Query Match 37.5%; Score 12; DB 9; Length 34;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 8 TTTTGTGCTTTT 19
|||||
15 TTTTGTGCTTTT 4

Db 15 TTTTGTGCTTTT 4

RESULT 5
BH863768 47 bp DNA linear GSS 05-AUG-2002
LOCUS SALK_094542 Arabidopsis thaliana T-DNA insertion lines Arabidopsis
DEFINITION thaliana genomic clone SALK_094542, genomic survey sequence.

ACCESSION BH863768
VERSION BH863768.1 GI:22099666
KEYWORDS GSS.
SOURCE Arabidopsis thaliana (thale cress)
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE 1 (bases 1 to 47)
Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R.,
Gadrihab, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L.,
Shim, P., Zimmerman, J. and Ecker, J.R.
A Sequence-Indexed Library of Insertion Mutations in the
Arabidopsis Genome
Unpublished (2001)
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of
T-DNA. This sequence lies within 300 bases of the 5' end of
AT5g37350.
Class: T-DNA tagged

FEATURES
source
1..47
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/ecotype="Col-0"
/db_xref="taxon:3702"
/clone_1ib="SALK_094542"
/note="PCR was performed on Arabidopsis thaliana lines
each of which contains one or more T-DNA insertion
elements. The resultant fragment for each line was
directly sequenced to determine the genomic sequence at
the site of insertion. Details of the protocols used can
be found at http://signal.salk.edu/cdna_protocols.html"

ORIGIN
Query Match 37.5%; Score 12; DB 8; Length 47;
Best Local Similarity 100.0%; Pred. No. 4.7e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CGTGTGTTTGTG 13
|||||
3 CGTGTGTTTGTG 14

Db 3 CGTGTGTTTGTG 14

RESULT 6
AU256787 49 bp mRNA linear EST 25-APR-2002
LOCUS AU256787 3'-directed mouse cDNA library Mus musculus cDNA clone
DEFINITION BED0008976 3', mRNA sequence.
ACCESSION AU256787
VERSION AU256787
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 49)
Kato, K. and Matoba, R.
Generation of expressed sequence tags from mouse brain
Unpublished (2002)
Contact: Kikuya Kato
Graduate School of Biological Sciences
Nara Institute of Science and Technology
8916-5 Takayama, Ikoma, Nara 630-0101, Japan
Tel: 81-743-72-5581
Fax: 81-743-72-5589

Email: kkarco@bs.aist-nara.ac.jp,
URL: http://love2.aist-nara.ac.jp/BEI/index.html.
Location/Qualifiers

FEATURES

source

ORIGIN

/organism="Mus musculus"
/mol_type="mRNA"
/db_xref="taxon:10090"
/clone="BED0008976"
/issue_type="brain"
/clone_lib="3'-directed mouse cDNA library"

Query Match

Best Local Similarity 37.5%; Score 12; DB 1; Length 49;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy

7 TTTTGTGCTTTT 18
36 TTTTGTGCTTTT 25

Db

9 TTTTGTGCTTTT 20
12 TTTTGTGCTTTT 23

RESULT 7

AZ472468 50 bp DNA linear GSS 04-OCT-2000
LOCUS 1M0287P08R Mouse 10kb plasmid UGCLM library Mus musculus genomic

DEFINITION

clone UGCLM0287P08 R, genomic survey sequence.

ACCESSION

VERSION

AZ472468.1 GI:10630593

KEYWORDS

GSS.

SOURCE

Mus musculus (house mouse)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE

1 (bases 1 to 50)
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

AUTHORS

Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
Isilan, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, T.,
Reilly, M., Rose, R., Stokes, R., Tingey, A., von
Niederhausern, A. and Wright, D., Weiss, R.

TITLE

Mouse whole genome scaffolding with paired end reads from 10kb
plasmid inserts

JOURNAL

Unpublished (2000)

COMMENT

Contact: Robert B. Weiss
University of Utah Genome Center
Rm 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
84112, USA

Tel: 801 585 5606
Fax: 801 585 7177

Email: ddunn@genetics.utah.edu

Insert Length: 10000 Std Error: 0.00

Plate: 0287 row: P column: 08

Seq primer: CACACAGAAACACGCTATGACC

Class: plasmid ends

High quality sequence stop: 50.

Location/Qualifiers

1. 50

/organism="Mus musculus"
/mol_type="genomic DNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="UGCLM0287P08"
/sex="Male"

/lab_host="E. coli strain XL10-Gold, TI-resistant, F-"
/clone_lib="Mouse 10kb plasmid UGCLM library"

/note="Vector: PWD42ny, Purified genomic DNA from M.
musculus C57BL/6J (male) was obtained from the Jackson
Laboratory Mouse DNA Resource
(http://www.jax.org/resources/documents/dnares/). The DNA
was hydrodynamically sheared by repeated passage through a
0.005 inch orifice at constant velocity. The sheared DNA
was blunt end-repaired with T4 DNA polymerase and T4
polynucleotide kinase. Adaptor oligonucleotides were
ligated to the blunt ends in high molar excess. The

Location/Qualifiers

1. 56

/organism="Arabidopsis thaliana"

/mol_type="genomic DNA"

/strain="Col-0"

/db_xref="taxon:3711"

/clone="GABI-Kat"

/sex="Male"

/lab_host="E. coli strain XL10-Gold, TI-resistant, F-"

/clone_lib="Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for
flanking sequence tag-based reverse genetics

Plant Mol. Biol. 53 (1-2), 247-259 (2003)

14756321

Strizhov, N., Li, Y., Rosso, M.G., Viehovever, P., Dekker, K.A. and
Weishaar, B.

High-throughput generation of sequence indexes from T-DNA
mutagenized Arabidopsis thaliana lines

Biotechniques 35 (6), 1164-1168 (2003)

4 (bases 1 to 56)

Rosso, M.G., Li, Y., Strizhov, N. and Weishaar, B.

Submitted (31-MAR-2004) Weishaar, B., Max-Planck-Institut fuer
Zoechseforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany

77015. Details on an insertion within the locus defined by BAC clone
sequence are described in References 1-3. The sequences are
generated at the MPI for Plant Breeding Research in the context of
the GABI-Kat project. GABI-Kat is part of the German plant genomics
program designated 'GABI'. Information on line availability can be
found at: http://www.mpiz-koeln.mpg.de/GABI-Kat/.

FEATURES

source

adapted DNA was purified and size-selected for a 9.5 to
10.5 kb range using preparative agarose gel
electrophoresis. Vector DNA was prepared from a derivative
of PWD42 (gi|4732114|gb|AF129072.1), a copy-number
inducible derivative of plasmid R1. The vector was ligated
with adaptors complementary to the insert adaptors and
purified. The sheared, adapted mouse DNA was annealed to
adapted vector DNA, and transformed into
chemically-competent E. coli XL10-Gold (Stratagene) cells
and selected for ampicillin resistance."

ORIGIN

Query Match
Best Local Similarity 37.5%; Score 12; DB 8; Length 50;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy

9 TTTTGTGCTTTT 20
12 TTTTGTGCTTTT 23

Db

RESULT 8
BX286679 56 bp DNA linear GSS 02-APR-2004
LOCUS Arabidopsis thaliana T-DNA flanking sequence GK-392H12-018271,
genomic survey sequence.

DEFINITION

Arabidopsis thaliana T-DNA flanking sequence GK-392H12-018271,
genomic survey sequence.

ACCESSION

VERSION

BX286679.1 GI:28885675

KEYWORDS

GSS.

SOURCE

Arabidopsis thaliana (thale cress)

ORGANISM

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.

REFERENCE

1
Li, Y., Rosso, M.G., Strizhov, N., Viehovever, P. and Weishaar, B.
The identification of T-DNA insertion mutants in Arabidopsis
thaliana

AUTHORS

Li, Y., Rosso, M.G., Strizhov, N., Viehovever, P. and Weishaar, B.
The identification of T-DNA insertion mutants in Arabidopsis
thaliana

TITLE

Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for
flanking sequence tag-based reverse genetics

JOURNAL

Plant Mol. Biol. 53 (1-2), 247-259 (2003)

MEDLINE

22755829

PUBMED

12874060

REFERENCES

2
Rosso, M.G., Li, Y., Strizhov, N., Reis, B., Dekker, K. and
Weishaar, B.

AUTHORS

Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for
flanking sequence tag-based reverse genetics

TITLE

Arabidopsis thaliana T-DNA mutagenized population (GABI-Kat) for
flanking sequence tag-based reverse genetics

JOURNAL

Plant Mol. Biol. 53 (1-2), 247-259 (2003)

MEDLINE

14756321

PUBMED

3

REFERENCES

Strizhov, N., Li, Y., Rosso, M.G., Viehovever, P., Dekker, K.A. and
Weishaar, B.

AUTHORS

High-throughput generation of sequence indexes from T-DNA
mutagenized Arabidopsis thaliana lines

TITLE

Biotechniques 35 (6), 1164-1168 (2003)

JOURNAL

4 (bases 1 to 56)

MEDLINE

14682050

PUBMED

4

REFERENCES

Rosso, M.G., Li, Y., Strizhov, N. and Weishaar, B.

AUTHORS

Submitted (31-MAR-2004) Weishaar, B., Max-Planck-Institut fuer
Zoechseforschung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany

TITLE

77015. Details on an insertion within the locus defined by BAC clone
sequence are described in References 1-3. The sequences are
generated at the MPI for Plant Breeding Research in the context of
the GABI-Kat project. GABI-Kat is part of the German plant genomics
program designated 'GABI'. Information on line availability can be
found at: http://www.mpiz-koeln.mpg.de/GABI-Kat/.

JOURNAL

Location/Qualifiers

COMMENT

1. 56
/organism="Arabidopsis thaliana"

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/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-392H12-018271"
/clone_1lb="Arabidopsis thaliana T-DNA insertion lines"
/ecotype="Col-0"
/note="PCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector pACT161 (Genbank accession number: AJ537514). The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. T-DNA derived sequences were removed."

```

ORIGIN

```

Query Match      37.5%; Score 12; DB 9; Length 56;
Best Local Similarity 100.0%; Pred. No. 4.6e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Oy      6 GTTTGTCGTTT 17
         |||||
Db      42 GTTTGTCGTTT 53

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```

RESULT 9
AL754515      57 bp      DNA      linear      GSS 01-APR-2004
LOCUS
DEFINITION
Arabidopsis thaliana T-DNA flanking sequence GK-054F09-012480,
genomic survey sequence.
AL754515
AL754515.1 GI:21487013
GSS

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```

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Arabidopsis thaliana (thale cress)

```

```

REFERENCE
AUTHORS
TITLE
Li, Y., Rosso, M.G., Strizhov, N., Viehoveer, P. and Weishaar, B.
GABI-Kat SimpleSearch: a flanking sequence tag (FST) database for
the identification of T-DNA insertion mutants in Arabidopsis
thaliana
Bioinformatics 19 (11), 1441-1442 (2003)

```

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JOURNAL
MEDLINE
PUBMED
22755829
12874060

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REFERENCE
AUTHORS
Rosso, M.G., Li, Y., Strizhov, N., Reiss, B., Dekker, K. and
Weishaar, B.

```

```

TITLE
An Arabidopsis thaliana T-DNA mutagenised population (GABI-Kat) for
flanking sequence tag-based reverse genetics
Plant Mol. Biol. 53 (1-2), 247-259 (2003)
23117147
14756321

```

```

JOURNAL
MEDLINE
PUBMED
14682050

```

```

REFERENCE
AUTHORS
TITLE
Strizhov, N., Li, Y., Rosso, M.G., Viehoveer, P., Dekker, K.A. and
Weishaar, B.
High-throughput generation of sequence indexes from T-DNA
mutagenised Arabidopsis thaliana lines
Biotechniques 35 (6), 1164-1168 (2003)

```

```

JOURNAL
PUBMED
14682050
4 (bases 1 to 57)
Rosso, M.G., Li, Y., Strizhov, N. and Weishaar, B.
Direct Subnucleon
Submitted (31-MAR-2004) Weishaar B., Max-Planck-Institut fuer
Zuechtungsforchung, Carl-von-Linne-Weg 10, Koeln, 50829, Germany.
This sequence has been recovered from the left border of the T-DNA.
It indicates an insertion within the locus defined by BAC clone
fi913. Details on the protocols used for generation of the sequence
are described in References 1-3. The sequences are generated at the
MPI for Plant Breeding Research in the context of the GABI-Kat
project. GABI-Kat is part of the German Plant Genomics program
designated 'GABI'. Information on line availability can be found
at: http://www.mpiz-koeln.mpg.de/GABI-Kat/.

```

COMMENT

FEATURES source

```

Location/Qualifiers
1..57
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/strain="Columbia 0"
/db_xref="taxon:3702"
/clone="GK-054F09-012480"
/clone_1lb="Arabidopsis thaliana T-DNA insertion lines"
/ecotype="Col-0"
/note="PCR was performed on DNA from Arabidopsis thaliana plants (T1) which were transformed with the T-DNA from vector pACT161 (Genbank accession number: AJ537514). The lines contain one or more T-DNA insertions. The DNA fragment(s) resulting from the PCR were directly sequenced to determine the genomic sequence flanking the insertion. T-DNA derived sequences were removed."

```

ORIGIN

```

Query Match      37.5%; Score 12; DB 9; Length 57;
Best Local Similarity 100.0%; Pred. No. 4.6e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Oy      7 TTTTGTGCTTT 18
         |||||
Db      46 TTTTGTGCTTT 57

```

```

RESULT 10
AV964458      60 bp      mRNA      linear      EST 14-MAR-2002
LOCUS
DEFINITION
AV964458 Nori Satoh unpublished cDNA library, larva Clona
interstitial cDNA clone c1v13e18 5', mRNA sequence.
AV964458
AV964458.1 GI:19454154
EST.

```

```

ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Clona intestinalis

```

```

REFERENCE
AUTHORS
TITLE
Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;
Phlebobranchia; Clonidae; Clona.
1 (bases 1 to 60)
Satoh, N., Satou, Y., Kohara, Y. and Shin-i, T.
Expressed genes in Clona intestinalis
Unpublished (2000)

```

COMMENT

```

Contact: Nori Satoh
Department of Zoology
Kyoto University
Sakyo-ku, Kyoto, Kyoto 606-8502, Japan
Tel: 81-75-753-4081
Fax: 81-75-705-1113
Email: satoh@abcidian.zool.kyoto-u.ac.jp.
Location/Qualifiers
1..60
/organism="Clona intestinalis"
/mol_type="mRNA"
/db_xref="taxon:7719"
/clone="c1v13e18"
/tissue_type="whole animal"
/dev stage="larva"
/clone_1lb="Nori Satoh unpublished cDNA library, larva"

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FEATURES source

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Location/Qualifiers
1..60
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/mol_type="mRNA"
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/clone="c1v13e18"
/tissue_type="whole animal"
/dev stage="larva"
/clone_1lb="Nori Satoh unpublished cDNA library, larva"

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ORIGIN

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Query Match      37.5%; Score 12; DB 2; Length 60;
Best Local Similarity 100.0%; Pred. No. 4.6e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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```

Oy      6 GTTTTGTGCTTT 17
         |||||
Db      14 GTTTTGTGCTTT 3

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```

RESULT 11
BH863639      62 bp      DNA      linear      GSS 05-AUG-2002
LOCUS
BH863639

```

DEFINITION SALK 094273 Arabidopsis thaliana TDNA insertion lines Arabidopsis thaliana genomic clone SALK_094273, genomic survey sequence.

ACCESSION BH63639

VERSION BH63639.1 GI:22099493

KEYWORDS GSS.

SOURCE ORGANISM Arabidopsis thaliana (thale cress)

Arabidopsis thaliana

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsids.

REFERENCE 1 (bases 1 to 62)

Alonso, J.M., Leisse, T.U., Barajas, P., Chen, H., Cheuk, R., Gadgil, C., Jeske, A., Karnes, M., Kim, C.U., Parker, H., Prednis, L., Shin, P., Zimmerman, J., and Ecker, J.R., 2001. A Sequence-Indexed Library of Insertion Mutations in the Arabidopsis Genome Unpublished (2001)

JOURNAL COMMENT Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGAL)
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu

FEATURES

source

1..62
location/Qualifiers
/organism="Arabidopsis thaliana"
/mol_type="genomic DNA"
/ecotype="Col-0"
/db_xref="taxon:3702"
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ORIGIN

Query Match 37.5%; Score 12; DB 8; Length 62;
Best Local Similarity 100.0%; Pred. No. 4.6e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 CGTCGTTTTC 13
18 CGTCGTTTTC 29

Db

RESULT 12
CB064048

LOCUS

DEFINITION PY03C06.y1 Haemochus contortus whole worm pAMP1 v1 Haemochus contortus cDNA 5', mRNA sequence.

ACCESSION CB064048

VERSION CB064048.1 GI:27809626

KEYWORDS EST.

SOURCE ORGANISM Haemochus contortus

Haemochus contortus

Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylida; Trichostrongyloidea; Haemochidae; Haemochinae; Haemochus.

REFERENCE 1 (bases 1 to 63)

Wyllie, T., Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Tsagaris, V., Gibbons, M., Ritzer, E., Bennett, J., Franklin, C., Tsagaris, V., Ronko, I., Kennedy, S., Maguire, J., Beck, C., Underwood, K., Steptoe, M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and Wilson, R.

TITLE The Washington Univ. Nematode EST Project, 1999

JOURNAL Unpublished (1999)

COMMENT Contact: McCarter JP
The Washington Univ. Nematode EST Project, 1999
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.edu

FEATURES

source

1..63
location/Qualifiers
/organism="Haemochus contortus"
/mol_type="mRNA"
/db_xref="taxon:6289"
/issue_type="whole organism"
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/note="Vector: PAMPI; Site 1: NotI; Site 2: SalI; The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. The cDNA was library using a modified oligo-dT priming (Dyna). PCR based Synthesis kit from Clontech. Directionally cloned into the UDG sites of PAMPI. Whole worm RNA was provided by Dr. Douglas Jansner of Washington State University (djaansner@vetmed.wsu.edu). Seq primer: Primer name ambiguous.

ORIGIN

Query Match 37.5%; Score 12; DB 6; Length 63;
Best Local Similarity 100.0%; Pred. No. 4.6e+03;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTTC 12
16 TCGTCGTTTTC 27

Db

RESULT 13
CB192029

LOCUS

DEFINITION PY23h12.y1 Haemochus contortus whole worm pAMP1 v1 Haemochus contortus cDNA 5', mRNA sequence.

ACCESSION CB192029

VERSION CB192029.1 GI:28255421

KEYWORDS EST.

SOURCE ORGANISM Haemochus contortus

Haemochus contortus

Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylida; Trichostrongyloidea; Haemochidae; Haemochinae; Haemochus.

REFERENCE 1 (bases 1 to 63)

Wyllie, T., Dante, M., Marra, M., Hillier, L., Kucaba, T., Theising, B., Tsagaris, V., Gibbons, M., Ritzer, E., Bennett, J., Franklin, C., Underwood, K., Steptoe, M., Allen, M., Person, B., Swaller, T., Harvey, N., Schurk, R., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCann, R., Waterston, R. and Wilson, R.

TITLE The Washington Univ. Nematode EST Project, 1999

JOURNAL Unpublished (1999)

COMMENT Contact: McCarter JP
The Washington Univ. Nematode EST Project, 1999
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 286 1800
Fax: 314 286 1810

Email: east@wustl.edu
The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dyna1). PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UDG sites of PAMPI. Whole worm RNA was provided by Dr. Douglas Jassmer of Washington State University (djasmer@vetmed.wsu.edu).
Seq primer: primer name ambiguous.

FEATURES

source

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Location/Qualifiers
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/clone_lib="Haemochus contortus whole worm PAMPI v1"
/note="Vector: PAMPI; Site 1: NotI; Site 2: SalI; The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dyna1). PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UDG sites of PAMPI. Whole worm RNA was provided by Dr. Douglas Jassmer of Washington State University (djasmer@vetmed.wsu.edu)."

ORIGIN

Query Match

Best Local Similarity 37.5%; Score 12; DB 6; Length 63;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGT 12
15 TCGTCGTTTGT 26

RESULT 14

CB191498

LOCUS

DEFINITION PY32h10.v1 Haemochus contortus whole worm PAMPI v1 Haemochus contortus cDNA 5', mRNA sequence.

ACCESSION

CB191498.1 GI:28254890

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

64 bp mRNA linear EST 05-FEB-2003
CB191498
PY32h10.v1 Haemochus contortus whole worm PAMPI v1 Haemochus contortus cDNA 5', mRNA sequence.
CB191498.1 GI:28254890
EST.
Haemochus contortus
Haemochus contortus
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Strongylidae; Trichostrongylidae; Haemonchidae; Haemonchinae; Haemochus.
1 (bases 1 to 64)
McCarter, J., Clifton, S., Chapell, B., Pape, D., Martin, J., Wylie, T., Dante, M., Maria, M., Hillier, L., Kucaba, T., Theising, B., Bowers, Y., Gibbons, M., Ritter, E., Bennett, J., Franklin, C., Tsagarashvili, R., Ronko, I., Kennedy, S., Maguire, L., Beck, C., Underwood, K., Stepcio, M., Allen, M., Person, B., Schaller, T., Harvey, N., Schurk, R., Kohn, S., Shin, T., Jackson, Y., Cardenas, M., McCan, R., Waterston, R. and Wilson, R.
The Washington Univ. Nematode EST Project, 1999
Unpublished (1999)
Contact: McCarter JP
The Washington Univ. Nematode EST Project, 1999
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108, USA
Tel: 314 266 1800
Fax: 314 266 1810
Email: east@wustl.edu
The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dyna1). PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UDG sites of PAMPI. Whole worm RNA was provided by Dr. Douglas Jassmer of Washington State University (djasmer@vetmed.wsu.edu).

FEATURES

source

Seq primer: primer name ambiguous.

Location/Qualifiers

1. .64

/organism="Haemochus contortus"

/mol_type="mRNA"

/db_xref="taxon:6289"

/tissue_type="whole organism"

/lab_host="DH10B"

/clone_lib="Haemochus contortus whole worm PAMPI v1"

/note="Vector: PAMPI; Site 1: NotI; Site 2: SalI; The library was constructed by Claire Murphy and Dr. James McCarter at Washington University, St. Louis. The cDNA was made by using Dynabead oligo-dT priming (Dyna1). PCR based library using a modified protocol from the SMART PCR cDNA Synthesis Kit from Clontech. Directionally cloned into the UDG sites of PAMPI. Whole worm RNA was provided by Dr. Douglas Jassmer of Washington State University (djasmer@vetmed.wsu.edu)."

ORIGIN

Query Match

Best Local Similarity 37.5%; Score 12; DB 6; Length 64;
Matches 12; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TCGTCGTTTGT 12
16 TCGTCGTTTGT 27

RESULT 15

BH848405

LOCUS

DEFINITION SALK_068149.48.55.x Arabidopsis thaliana TDNA insertion lines Arabidopsis thaliana genomic clone SALK_068149.48.55.x, genomic survey sequence.

ACCESSION

BH848405

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

23 bp DNA linear GSS 13-JUN-2002
BH848405
SALK_068149.48.55.x Arabidopsis thaliana TDNA insertion lines Arabidopsis thaliana genomic clone SALK_068149.48.55.x, genomic survey sequence.
BH848405
BH848405.1 GI:21419276
GSS.
Arabidopsis thaliana (thale cress)
Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
1 (bases 1 to 23)
Alonso, J.M., Leisse, T.J., Barajas, P., Chen, H., Cheuk, R., Gadrinab, C., Jeske, A., Karnes, M., Kim, C.J., Parker, H., Prednis, L., Shin, P., Zimmermann, J. and Ecker, J.R.
A Sequence-indexed Library of Insertion Mutations in the Arabidopsis Genome
Unpublished (2001)
Contact: Joseph R. Ecker
Salk Institute Genomic Analysis Laboratory (SIGAL)
The Salk Institute for Biological Studies
10010 N. Torrey Pines Road, La Jolla, CA 92037, USA
Tel: 858 453 4100 x1752
Fax: 858 558 6379
Email: ecker@salk.edu
This is single pass sequence recovered from the left border of TDNA.
Class: TDNA tagged.
Location/Qualifiers
1. .23
/organism="Arabidopsis thaliana"
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/note="PCR was performed on Arabidopsis thaliana lines each of which contains one or more TDNA insertion elements. The resultant fragment for each line was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can

ORIGIN be found at http://signal.salk.edu/tdna_protocols.html

Query Match 34.4%; Score 11; DB 8; Length 23;
 Best local similarity 100.0%; Pred. No. 1.9e+04;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 TCGTCGTTTG 11
 |||||
 Db 6 TCGTCGTTTG 16

Search completed: February 14, 2005, 09:29:29
 Job time : 7364 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM protein - protein search, using sw model

Run on: February 8, 2005, 17:14:08 / Search time 170 Seconds
(without alignments)
102.378 Million cell updates/sec

Title: US-10-076-674A-9
Perfect score: 243
Sequence: 1 TAKSKRPPSTATYQFGGLS.....IVHRLGVGGEHNSYGLRPG 45

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2105692 seqs, 386760381 residues
Total number of hits satisfying chosen parameters: 2105692

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq.16Dec04:*
1: geneseqp1980s:*
2: geneseqp1990s:*
3: geneseqp2000s:*
4: geneseqp2001s:*
5: geneseqp2002s:*
6: geneseqp2003as:*
7: geneseqp2003bs:*
8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	243	100.0	45	2	AAR62721
2	243	100.0	45	7	ADD89949 LHRH-cont
3	243	100.0	45	8	ADJ56908 Human LHR
4	235	96.7	45	3	AAY68573 Peptide 1
5	235	96.7	45	3	AAY91165 Modified
6	216.5	89.1	46	2	AAR62728 LHRH-cont
7	213	87.7	49	3	AAY91177 Modified
8	211	86.8	45	3	AAY68577 Peptide 1
9	211	86.8	45	3	AAY91172 Modified
10	208	85.6	47	3	AAY68583 Peptide 1
11	208	85.6	47	3	AAY91180 Inv epit
12	206	84.8	45	3	AAY91166 Modified
13	203	83.5	47	3	AAY68586 Peptide 1
14	203	83.5	47	3	AAY91183 Inv epit
15	199	81.9	45	3	AAY68571 Peptide 1
16	187	77.0	49	3	AAY68585 Peptide 1
17	187	77.0	49	3	AAY91178 Modified
18	187	77.0	49	3	AAY91182 Inv epit
19	181	74.5	49	3	AAY68580 Peptide 1
20	180	74.1	35	2	AAR65381 Universal
21	169.5	69.8	48	2	AAR62725 LHRH-cont
22	159	65.4	49	2	AAR62724 LHRH-cont
23	157	64.6	45	7	ADD89948 LHRH-pept
24	157	64.6	45	8	ADJ56907 Human LHR
25	156.5	64.4	54	2	AAR62722 LHRH-cont

26	156	64.2	45	2	AAR62720	AAR62720 LHRH-cont
27	152	62.6	47	2	AAR62723	AAR62723 LHRH-cont
28	151.5	62.3	46	3	AAY68595	AAY68595 Peptide 1
29	151.5	62.3	46	3	AAY91195	AAY91195 Inv epit
30	150.5	61.9	48	2	AAR62729	AAR62729 LHRH-cont
31	150	61.7	42	2	AAR62708	AAR62708 LHRH-cont
32	150	61.7	80	3	AAY68530	AAY68530 Synthetic
33	147	60.5	27	2	AAR62707	AAR62707 LHRH-cont
34	147	60.5	27	3	AAY68567	AAY68567 Peptide 1
35	147	60.5	27	3	AAY91156	AAY91156 MVF Th ep
36	144	59.3	27	3	AAY91163	AAY91163 Modified
37	139	57.2	27	3	AAY91161	AAY91161 Modified
38	139	57.2	27	3	AAY91167	AAY91167 Modified
39	137	56.4	28	3	AAY91158	AAY91158 Modified
40	134	55.1	27	3	AAY68575	AAY68575 Peptide 1
41	134	55.1	27	3	AAY91170	AAY91170 Modified
42	133	54.7	31	3	AAY91175	AAY91175 Modified
43	131	53.9	28	2	AAR62726	AAR62726 LHRH-cont
44	130	53.5	28	3	AAY91159	AAY91159 Modified
45	130	53.5	31	3	AAY91179	AAY91179 Modified

ALIGNMENTS

RESULT 1	
AAR62721	AAR62721 standard; peptide; 45 AA.
XX	
AC	AAR62721;
XX	
DT	25-MAR-2003 (revised)
DT	10-SEP-1995 (first entry)
XX	
DE	LHRH-containing immunogenic peptide.
XX	
KW	Helper T cell epitope; universal immune stimulator; invasive; hapten;
KW	vaccine; LHRH; luteinising hormone releasing hormone; prostatic;
KW	androgen-dependent carcinoma; antitumour; infertility;
KW	measles virus F protein.
XX	
OS	Synthetic.
XX	
FH	Key
FT	Domain
FT	Domain
FT	Domain
FT	Domain
FT	Domain
XX	
PD	10-NOV-1994.
XX	
PE	28-APR-1994; 94MO-US004832.
XX	
PR	27-APR-1993; 93US-00057166.
PR	14-APR-1994; 94US-00229275.
XX	
PA	(LADD/) LADD A E.
PA	(WANG/) WANG C Y.
PA	(ZAMB/) ZAMB T.
XX	
PI	Ladd AE, Wang CY, Zamb T;
DR	WPI; 1994-357910/44.
XX	
PT	Immunogenic luteinising hormone releasing hormone peptide(s) - that
PT	suppress LHRH activity in males and females.
XX	
PS	Claim 8; Page 88; 213pp; English.

CC Synthetic immunogenic peptides are provided in which a universal immune
CC stimulator is linked to a peptide or protein hapten containing B cell
CC and/or cytotoxic T lymphocyte epitopes, giving a product which causes
CC potent immune responses to the coupled peptide or protein. The stimulator
CC consists of (A) a promiscuous helper T cell epitope (TH) which elicits an
CC immune response to the coupled peptide in members of a heterogeneous
CC population expressing diverse HLA phenotypes, and (B) an adjuvant peptide
CC sequence from the invasive protein of Yersinia. Spacer amino acid
CC sequences (e.g. Gly-Gly) can be provided between the invasive and TH
CC domains and between the immune stimulator and hapten components. When the
CC hapten is LHRH, then optionally the invasive domain can be omitted from
CC the immune stimulator component. The present sequence represents an LHRH-
CC containing immunogenic peptide as above which can be used as a potent
CC vaccine for treating e.g. prostatic hyperplasia, androgen-dependent
CC carcinoma, prostatic carcinoma, testicular carcinoma, endometriosis,
CC benign uterine tumors, recurrent functional ovarian cysts, (severe)
CC premenstrual syndrome or oestrogen-dependent breast cancer, or for
CC induction of infertility. (updated on 25-MAR-2003 to correct FN field.)
SQ Sequence 45 AA;

Query Match 100.0%; Score 243; DB 2; Length 45;
Best Local Similarity 100.0%; Pred. No. 4.1e-27;
Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPPSYATYQFGSLSEIKGYVHRLEGVGGEHMSYGLRPG 45
DB 1 TAKSKKPPSYATYQFGSLSEIKGYVHRLEGVGGEHMSYGLRPG 45

RESULT 2
ADD89949 ID ADD89949 standard; protein; 45 AA.
AC ADD89949;

DT 29-JAN-2004 (first entry)

DE LHRH peptide used in immunostimulant complex for prostate cancer vaccine.

XX Immunostimulant; vaccine; human; immunogen; LHRH; immunotherapy;

KW prostate cancer.

OS Synthetic.

OS Homo sapiens.

PN MO2003068169-A2.

PD 21-AUG-2003.

PF 14-FEB-2003; 2003WO-US004711.

PR 14-FEB-2002; 2002US-00076674.

PR 31-JAN-2003; 2003US-00076674.

PA (UNBI-) UNITED BIOMEDICAL INC.

PI Sokoll KK;

WPI; 2003-778890/73.

PT Stabilized immunostimulating complex, useful for vaccination, e.g.
PT against human immune deficiency viruses, comprises cationic peptide
PT immunogen and anionic oligonucleotide.

PS Claim 17; SEQ ID NO 9; 159bp; English.

CC The present sequence is that of a synthetic immunogenic peptide derived
CC from human LHRH. This is an example of peptides that can be used in
CC claimed immunostimulatory complexes of the invention that are
CC specifically adapted to act as adjuvant and as peptide immunogen
CC stabiliser. The complexes comprise a Cpg oligonucleotide and a
CC biologically active peptide immunogen. The complex is particulate and can

CC efficiently present peptide immunogens to the cells of the immune system
CC to produce an immune response. The complexes may be prepared with various
CC ratios of peptides to Cpg oligonucleotides to provide different physical
CC properties, such as the size of the microparticle. An immunostimulatory
CC complex comprising the present LHRH derived peptide can be used in a
CC vaccine for prostate cancer.
SQ Sequence 45 AA;

Query Match 100.0%; Score 243; DB 7; Length 45;
Best Local Similarity 100.0%; Pred. No. 4.1e-27;
Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPPSYATYQFGSLSEIKGYVHRLEGVGGEHMSYGLRPG 45
DB 1 TAKSKKPPSYATYQFGSLSEIKGYVHRLEGVGGEHMSYGLRPG 45

RESULT 3

ID ADJ56908 standard; peptide; 45 AA.

AC ADJ56908;

DT 06-MAY-2004 (first entry)

DE Human LHRH immunogenic peptide #3.

KW Immunostimulatory complex; adjuvant; peptide immunogen stabiliser;

KW water-in-oil emulsion; suspension; vaccine; prostate cancer;

KW hormone ablation; allergy; HIV infection; foot-and-mouth disease;

KW therapy; human; antigen; LHRH.

OS Homo sapiens.

PN US2004009897-A1.

PD 15-JAN-2004.

PF 21-MAY-2003; 2003US-00355161.

PR 14-FEB-2002; 2002US-00076674.

PA (SOKO/) SOKOLL K K.

PI Sokoll KK;

WPI; 2004-212745/20.

PT Stabilized immunostimulatory complex useful for treating allergy, HIV
PT infection or prostate cancer, comprising cationic peptide immunogen and
PT anionic Cpg oligonucleotide.

PS Claim 17; SEQ ID NO 9; 63bp; English.

CC The invention relates to an immunostimulatory complex specifically
CC adapted to act as adjuvant and as a peptide immunogen stabiliser. The
CC invention is useful for preparing a water-in-oil emulsion, suspension and
CC vaccine. It is also useful for treating prostate cancer, hormone
CC ablation, allergy, HIV infection, foot-and-mouth disease, etc. The
CC present sequence is human LHRH immunogenic peptide used in the invention.
SQ Sequence 45 AA;

Query Match 100.0%; Score 243; DB 8; Length 45;
Best Local Similarity 100.0%; Pred. No. 4.1e-27;
Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPPSYATYQFGSLSEIKGYVHRLEGVGGEHMSYGLRPG 45
DB 1 TAKSKKPPSYATYQFGSLSEIKGYVHRLEGVGGEHMSYGLRPG 45

Sequence	AA	Sequence	AA
1	AAV68573	1	AAV68573
2	AAV68573 standard; peptide; 45 AA.	2	AAV68573
3	05-MAY-2000 (first entry)	3	Peptide immunogen comprising a Th epitope and LHRH target antigen.
4	Peptide immunogen comprising a Th epitope and LHRH target antigen.	4	Helper T cell epitope; F protein; Measles virus; peptide immunogen; LHRH;
5	luteinizing hormone-releasing hormone; spermatogenesis; ovulation;	5	testosterone; sexual development; sex hormone; promiscuous T helper epitope;
6	vaccine; contraceptive; hormone-dependent tumour; prostate cancer;	6	breast cancer; endometriosi; boar taint; meat quality; invasive domain;
7	immunocastration.	7	Immunocastration.
8	Synthetic.	8	Synthetic.
9	Measles virus.	9	Measles virus.
10	Unidentified.	10	Unidentified.
11	Key	11	Location/Qualifiers
12	Peptide	12	1..16
13	Peptide	13	/note="invasive domain AAV68565"
14	Peptide	14	17..18
15	Peptide	15	/note="spacer"
16	Peptide	16	19..33
17	Peptide	17	/note="helper Th epitope AAV68544"
18	Peptide	18	34..35
19	Peptide	19	/note="spacer"
20	Peptide	20	36..45
21	Peptide	21	/note="LHRH antigenic epitope AAV68566"
22	Peptide	22	MO966952-AA.
23	Peptide	23	29-DEC-1999.
24	Peptide	24	21-JUN-1999; 99MO-US013960.
25	Peptide	25	20-JUN-1998; 98US-0010041.
26	Peptide	26	(UNBI-) UNITED BIOMEDICAL INC.
27	Peptide	27	Wang CY;
28	Peptide	28	WPI; 2000-160562/14.
29	Peptide	29	New peptide immunogen containing luteinizing hormone-releasing hormone
30	Peptide	30	antigen site and helper T cell epitope, for e.g. contraception and
31	Peptide	31	treatment of cancer.
32	Peptide	32	Claim 9; Page 71; 102pp; English.
33	Peptide	33	The present sequence represents a peptide immunogen comprising an invasive
34	Peptide	34	domain immunostimulatory peptide of Yersinia sp., a synthetic helper T
35	Peptide	35	cell (Th) epitope and a target antigen, luteinizing hormone-releasing
36	Peptide	36	hormone (LHRH). The synthetic Th epitope is derived from a structured
37	Peptide	37	synthetic antigen library (SSAL) designated SSAL Th. SSAL Th is
38	Peptide	38	modelled after a promiscuous epitope taken from the F protein of the
39	Peptide	39	Measles virus. The peptide immunogens cause induction of a specific
40	Peptide	40	immune response to LHRH which is involved in regulation of
41	Peptide	41	spermatogenesis, ovulation, oestrus, sexual development and secretion of
42	Peptide	42	sex hormones. Provision of a promiscuous T helper epitope (which is
43	Peptide	43	functional in genetically diverse subjects) provides optimum
44	Peptide	44	immunogenicity to the B cell epitopes of the target antigen and thus high
45	Peptide	45	antibody titres against the target antigen. The peptide immunogens of the
46	Peptide	46	invention are used to vaccinate against mammalian LHRH, for use as
47	Peptide	47	(reversible) contraceptive; control of hormone-dependent tumours (cancer
48	Peptide	48	of prostate or breast, also endometriosi); to prevent boar taint (and
49	Peptide	49	improve meat quality) and for immunocastration
50	Peptide	50	Sequence 45 AA;

Query Match	96.7%	Score 235	DB 3	Length 45
Best Local Similarity	91.1%	Pred. No. 5.7e-26		
Matches	41	Conservative	4	Mismatches 0; Indels 0; Gaps 0;
Qy	1	TAKSKKPEPYATATQFGLSBKIVYHRLGEGHMSYGLRQ	45	
Db	1	TAKSKKPEPYATATQFGLSBKIVYHRLGEGHMSYGLRQ	45	
RESULT 5				
ID	AA91165	standard; peptide; 45 AA.		
XX	AA91165;			
AC				
XX				
DT	12-SEP-2003	(revised)		
XX				
DT	22-MAY-2000	(first entry)		
XX				
DE	Modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:45.			
XX				
KM	Promiscuous T-cell epitope; measles virus F protein; MVF;			
KM	hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;			
KM	luteinizing hormone releasing hormone; LHRH; contraceptive; anticancer;			
KM	somatostatin; growth promotion; CD4 receptor; HIV-1; antiviral; PMDV;			
KM	foot and mouth disease virus; immunoglobulin E; IGE; anti-allergic;			
KM	Plasmodium falciparum; circumsporozoite; antimalarial; CESTP;			
KM	cholesterol ester transport protein; anti-arteriosclerotic.			
XX				
OS	Measles virus.			
OS	Rattus sp.			
OS	Chimeric.			
XX				
PN	W09966957-A2.			
XX				
PD	29-DEC-1999.			
XX				
PF	21-JUN-1999; 99WO-US013975.			
XX				
PR	20-JUN-1998; 98US-00100412.			
XX				
PA	(UNBI-) UNITED BIOMEDICAL INC.			
PI				
XX	Wang CY;			
DR				
XX	WPI; 2000-160564/14.			
XX				
CC	The invention relates to novel promiscuous T helper cell epitopes (Th),			
CC	and immunogenic peptides comprising the Th epitopes of the invention			
CC	along with B cell epitopes. The Th epitopes and peptide immunogens			
CC	containing them, are used to induce a T helper cell response,			
CC	specifically against Plasmodium falciparum, cholesterol ester transport			
CC	protein (CESTP) or HIV epitopes, but more generally against any pathogen,			
CC	immunoreactive self-antigen or tumour antigen. The Th epitopes and			
CC	peptide immunogens may be used for prevention and/or treatment of			
CC	infections (HIV, foot-and-mouth disease or malaria); for cancer			
CC	immunotherapy; for inhibition of the action of luteinizing hormone			
CC	releasing hormone (LHRH) for contraception, treatment of hormone-			
CC	dependent cancer, prevention of boar taint in meat, and immunosuppression			
CC	for promoting the growth of animals; or for treating allergies or			
CC	arteriosclerosis. Incorporation of a promiscuous Th (functional in			
CC	genetically diverse subjects) into an immunogen improves capacity to			
CC	induce a strong T helper cell-mediated immune response, resulting in			
CC	production of antibodies against a target antigen. Th can replace carrier			
CC	proteins and pathogen-derived T helper epitopes. Sequence AA91121			
CC	represents a promiscuous T helper epitope from the measles virus F (MVF)			
CC	protein and sequences AA91122-Y91142, AA911226 and AA911245-Y91246			

CC represent synthetic Th epitopes based on the MZF Th epitope. Sequence
 CC AAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV)
 CC surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes
 CC derived from this HBV epitope. AAY91156-Y91196, AAY91227 and AAY91242-
 CC Y91244 are antigenic peptides comprising an LHRH sequence joined to a
 CC promiscuous Th epitope. AAY91197 is the LHRH target antigenic peptide
 CC used in these LHRH antigenic peptides. AAY91200 is somatostatin, and
 CC AAY91201-Y91207 are antigenic peptides comprising somatostatin and a Th
 CC epitope. Somatostatin immunogens may be used to promote growth in
 CC livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and
 CC AAY91209-Y90211 are MZF Th epitope/CD4 CDR2 antigenic peptides which
 CC be used to prevent HIV infection of T cells. AAY90212 is a modified
 CC version of a human IGE (immunoglobulin E) CH3 domain, and AAY90213-Y90219
 CC are Th epitope/IGE CH3 antigenic peptides which may be used in the
 CC treatment of allergies. AAY91220 is a peptide derived from foot and mouth
 CC disease virus (FMDV) VP1 capsid protein and AAY91221-Y91222 comprise this
 CC peptide and a Th epitope. AAY91223 is a Plasmodium falciparum
 CC circumsporozoite (CS) target antigen, and AAY91224-Y91225 comprise this
 CC antigen and an MZF Th epitope, and may be used in a malaria vaccine.
 CC AAY91228-Y91231 represent CERP-derived peptides and AAY91232-Y91241
 CC immunogens comprising a CERP peptide and a Th epitope which may be used
 CC to prevent or treat arteriosclerosis and cardiovascular disease. AAY91247
 CC and AAY91252-Y91257 are HIV-1 neutralising B-cell epitopes, and AAY91248-
 CC Y91251 and AAY91258-Y91273 are antigenic peptides comprising MZF Th and
 CC HIV-1 B-cell epitope which may be used as a component in an anti-HIV-1
 CC vaccine. AAY91198 and AAY91199 are respectively an immunostimulatory
 CC invasive protein epitope from Yersinia species, and a hinge spacer peptide,
 CC both of which may optionally be used in the antigenic peptide.
 CC invention. (Updated on 12-SEP-2003 to standardise OS field)

XX Sequence 45 AA;

SO Query Match 96.7%; Score 235; DB 3; Length 45;
 Best Local Similarity 91.1%; Pred. No. 5.7e-26;
 Matches 41; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPPSYTATYQFGSLSEIKGIVHRLRGVGGHMSYGLRPG 45
 DB 1 TAKSKKPPSYTATYQFGSLSEIKGIVHRLRGVGGHMSYGLRPG 45

RESULT 6
 AAR62728 ID AAR62728 standard; peptide; 46 AA.
 AC AAR62728;

DT 25-MAR-2003 (revised)
 DT 17-SEP-1995 (first entry)

DE LHRH-containing immunogenic peptide.

KW Helper T cell epitope; universal immune stimulator; invasive; hapten;
 KW vaccine; LHRH, luteinising hormone releasing hormone; prostaglandin;
 KW androgen-dependent carcinoma; antitumour; infertility;
 KW structured synthetic antigen library; SSAL.

OS Synthetic.

XX Key Location/Qualifiers

XX Domain 1-16
 XX Domain /note= "invasin domain"

XX Domain 19-34
 XX /note= "structured synthetic antigen library (see US Ser.
 No. 143312, 26 Oct 1993), where the variant positions
 noted below may be a mixture of the specified residues.
 This domain functions as a helper T cell epitope"

XX Misc-difference 19 /label= Asp, Glu

XX Misc-difference 20 /label= Leu, Ile, Val, Phe

XX Misc-difference 22 /label= Glu, Asp

FT Misc-difference 23 /label= Leu, Ile, Val, Phe
 FT Misc-difference 24 /label= Lys, Arg
 FT Misc-difference 26 /label= Leu, Ile, Val, Phe
 FT Misc-difference 27 /label= Leu, Ile, Val, Phe
 FT Misc-difference 28 /label= Leu, Ile, Val, Phe
 FT Misc-difference 30 /label= Leu, Ile, Val, Phe
 FT Misc-difference 31 /label= Lys, Arg
 FT Misc-difference 32 /label= Leu, Ile, Val, Phe
 FT Misc-difference 34 /label= Glu, Asp
 FT Domain /label= Leu, Ile, Val, Phe
 FT /note= "LHRH hapten"

XX WO9425060-A1.

XX 10-NOV-1994.

XX 28-APR-1994; 94MO-US04832.

XX 27-APR-1993; 93US-00057166.

XX 14-APR-1994; 94US-00229275.

XX (LADD/) LADD A E.

XX (WANG/) WANG C Y.

XX (ZAMB/) ZAMB T.

XX Ladd AE, Wang CY, Zamb T;

XX WPI, 1994-357910/44.

XX Immunogenic luteinising hormone releasing hormone peptide(s) - that
 suppress LHRH activity in males and females.

PS Claim 8; Page 88; 213pp; English.

XX Synthetic immunogenic peptides are provided in which a universal immune
 CC stimulator is linked to a peptide or protein hapten containing B cell
 CC and/or cytotoxic T lymphocyte epitopes, giving a product which causes
 CC potent immune responses to the coupled peptide or protein. The stimulator
 CC consists of (A) a promiscuous helper T cell epitope (Th) which elicits an
 CC immune response to the coupled peptide in members of a heterogeneous
 CC population expressing diverse HLA phenotypes, and (B) an adjuvant peptide
 CC sequence from the invasive protein of Yersinia. Spacer amino acid Th
 CC domains and between the immune stimulator and hapten components. When the
 CC hapten is LHRH, then optionally the invasive domain can be omitted from
 CC the immune stimulator component. The present sequence represents an LHRH-
 CC containing immunogenic peptide as above in which the Th is a structured
 CC synthetic antigen library (SSAL). The peptide can be used as a potent
 CC vaccine for treating e.g. prostatic hyperplasia, androgen-dependent
 CC carcinoma, prostatic carcinoma, testicular carcinoma, endometriosis,
 CC benign uterine tumours, recurrent functional ovarian cysts, (beware)
 CC premenstrual syndrome or oestrogen-dependent breast cancer, or for
 CC induction of infertility. (Updated on 25-MAR-2003 to correct PN field.)

XX Sequence 46 AA;

SO Query Match 89.1%; Score 216.5; DB 2; Length 46;
 Best Local Similarity 84.8%; Pred. No. 2.7e-23;
 Matches 39; Conservative 6; Mismatches 0; Indels 1; Gaps 1;

QY 1 TAKSKKPPSYTATYQFGSLSEIKGIVHRLRGVGGHMSYGLRPG 45
 DB 1 TAKSKKPPSYTATYQFGSLSEIKGIVHRLRGVGGHMSYGLRPG 46

RESULT 7
 AAY91177
 ID AAY91177 standard; peptide: 49 AA.
 AC AAY91177;
 XX
 DT 12-SEP-2003 (revised)
 DT 22-MAY-2000 (first entry)
 DE Modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:57.
 XX
 KM Promiscuous T-cell epitope; measles virus F protein; MVF;
 KM hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;
 KM interleukin-6 releasing hormone; LHRH; contraceptive; anticancer;
 KM somatostatin; growth promotion; CD4 receptor; HIV-1; antiviral; FMDV;
 KM foot and mouth disease virus; immunoglobulin E; IGE; anti-allergic;
 KM Plasmodium falciparum; circumsporozoite; antimalarial; CERP;
 KM cholesterol ester transport protein; anti-arteriosclerotic.
 XX
 OS Measles virus.
 OS Rattus sp.
 OS Chimeric.
 XX
 MO9966957-A2.
 XX
 PD 29-DEC-1999.
 XX
 PF 21-JUN-1999; 99MO-US013975.
 XX
 PR 20-JUN-1998; 98US-00100412.
 XX
 PA (UNBI-) UNITED BIOMEDICAL INC.
 XX
 PI Wang CY;
 XX
 DR WPI; 2000-160564/14.
 XX
 PT New artificial T helper cell epitope and derived immunogens with target
 PT antigenic site, for immunization against e.g. malaria, arteriosclerosis
 PT or human immune deficiency virus.
 XX
 PS Example 1; Page 85; 129pp; English.
 XX
 CC The invention relates to novel promiscuous T helper cell epitopes (Th),
 CC and immunogenic peptides comprising the Th epitopes of the invention
 CC along with B cell epitopes. The Th epitopes and peptide immunogens
 CC containing them, are used to induce a T helper cell response,
 CC specifically against Plasmodium falciparum, cholesterol ester transport
 CC protein (CERP) or HIV epitopes, but more generally against any pathogen,
 CC immunoreactive self-antigen or tumour antigen. The Th epitopes and
 CC peptide immunogens may be used for prevention and/or treatment of
 CC infections (HIV, foot-and-mouth disease or malaria); for cancer
 CC immunotherapy; for inhibition of the action of interleukin hormone
 CC releasing hormone (LHRH) for contraception, treatment of hormone-
 CC dependent cancer, prevention of boar taint in meat, and immunocastration)
 CC ; for promoting the growth of animals; or for treating allergies or
 CC arteriosclerosis. Incorporation of a promiscuous Th (functional in
 CC genetically diverse subjects) into an immunogen improves capacity to
 CC induce a strong T helper cell-mediated immune response, resulting in
 CC production of antibodies against a target antigen. Th can replace carrier
 CC proteins and pathogen-derived T helper epitopes. Sequence AAY91121
 CC represents a promiscuous T helper epitope from the measles virus F (MVF)
 CC protein and sequences AAY91122-Y91142, AAY91126 and AAY91245-Y91246
 CC represent synthetic Th epitopes based on the MVF Th epitope. Sequence
 CC AAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV)
 CC surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes
 CC derived from this HBV epitope. AAY91156-Y91166, AAY91227 and AAY91242-
 CC Y91244 are antigenic peptides comprising an LHRH sequence joined to a
 CC promiscuous Th epitope. AAY91197 is the LHRH target antigenic peptide
 CC used in these LHRH antigenic peptides. AAY91200 is somatostatin, and
 CC AAY91201-Y91207 are antigenic peptides comprising somatostatin and a Th
 CC epitope. Somatostatin immunogens may be used to promote growth in

CC livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and
 CC AAY91209-Y90211 are MVA Th epitope/CD4 CDR2 antigenic peptides which may
 CC be used to prevent HIV infection of T cells. AAY90212 is a modified
 CC version of a human IGE (immunoglobulin E) CH3 domain, and AAY90213-Y90219
 CC are Th epitope/IGE CH3 antigenic peptides which may be used in the
 CC treatment of allergies. AAY91220 is a peptide derived from foot and mouth
 CC disease virus (FMDV) VP1 capsid protein and AAY91221-Y91222 comprise this
 CC peptide and a Th epitope. AAY91223 is a Plasmodium falciparum
 CC circumsporozoite (CS) target antigen, and AAY91224-Y91225 comprise the CS
 CC antigen and an MVA Th epitope and may be used in a malaria vaccine.
 CC AAY91228-Y91231 represent CERP-derived peptides and AAY91232-Y91241 are
 CC immunogens comprising a CERP peptide and a Th epitope which may be used
 CC to prevent or treat arteriosclerosis and cardiovascular disease. AAY91247
 CC and AAY91252-Y91257 are HIV-1 neutralising B-cell epitopes, and AAY91248-
 CC Y91251 and AAY91258-Y91273 are antigenic peptides comprising HIV Th and
 CC HIV-1 B-cell epitope which may be used as a component in an anti-HIV-1
 CC vaccine. AAY91198 and AAY91199 are respectively an immunostimulatory
 CC invasion protein epitope from Yersinia species, and hinge spacer peptide,
 CC both of which may optionally be used in the antigenic peptides of the
 CC invention. (Updated on 12-SEP-2003 to standardise OS field)
 CC
 XX
 SQ Sequence 49 AA;
 XX
 Query Match 87.7%; Score 213; DB 3; Length 49;
 Best Local Similarity 83.7%; Pred. No. 9.2e-23;
 Matches 41; Conservative 4; Mismatches 0; Indels 4; Gaps 2;
 XX
 QY 1 TASSKKPPSYATATYQFGG--LSEIKGVIVARLGGV--GGERHWSYGLRPG 45
 Db 1 TASSKKPPSYATATYQFGGISEIKGVIVARLGGVIVARLGGHWSYGLRPG 49
 XX
 RESULT 8
 AAY68577
 ID AAY68577 standard; peptide: 45 AA.
 XX
 AC AAY68577;
 XX
 DT 05-MAY-2000 (first entry)
 XX
 DE Peptide immunogen comprising a Th epitope and LHRH target antigen.
 XX
 KM Helper T cell epitope; F protein; Measles virus; peptide immunogen; LHRH;
 KM interleukin-6 releasing hormone; spermatogenesis; ovulation;
 KM oestrus; sexual development; sex hormone; promiscuous T helper epitope;
 KM vaccine; contraceptive; hormone-dependent tumour; prostate cancer;
 KM breast cancer; endometriosis; boar taint; meat quality; invasion domain;
 KM immunocastration.
 XX
 OS Synthetic.
 OS Yersinia sp.
 OS Measles virus.
 OS Undifferentiated.
 XX
 FH Key
 FT Peptide
 FT /note="invasion domain AAY68565"
 FT Peptide
 FT /note="spacer"
 FT Peptide
 FT /note="helper Th epitope AAY68547"
 FT Peptide
 FT /note="spacer"
 FT /note="LHRH antigenic epitope AAY68566"
 XX
 PN MO9966952-A1.
 XX
 PD 29-DEC-1999.
 XX
 PF 21-JUN-1999; 99MO-US013960.
 XX
 PR 20-JUN-1998; 98US-00100414.

XX (UNBI-) UNITED BIOMEDICAL INC.
 PA Wang CY;
 PI WPI, 2000-160562/14.
 DR WPI, 2000-160562/14.
 XX
 PT New peptide immunogen containing luteinizing hormone-releasing hormone
 PT antigen site and helper T cell epitope, for e.g. contraception and
 PS treatment of cancer.
 XX
 PS Claim 9; Page 73; 102pp; English.
 XX
 CC The present sequence represents a peptide immunogen comprising an invasin
 CC domain immunostimulatory peptide of *Yersinia sp.*, a synthetic helper T
 CC cell (Th) epitope and a target antigen, luteinizing hormone-releasing
 CC hormone (LHRH). The synthetic Th epitope is derived from a structured
 CC synthetic antigen library (SSAL) designated SSAL1 Th1. SSAL Th1 is
 CC modelled after a promiscuous epitope taken from the F protein of the
 CC Measles virus. The peptide immunogens cause induction of a specific
 CC immune response to LHRH which is involved in regulation of
 CC spermatogenesis, ovulation, oestrus, sexual development and secretion of
 CC sex hormones. Provision of a promiscuous T helper epitope (which is
 CC immunogenicity to the B cell epitopes of the target antigen and thus high
 CC antibody titres against the target antigen. The peptide immunogens of the
 CC (reversible) contraceptive, control of hormone-dependent tumours (cancer
 CC of prostate or breast, also endometriosis); to prevent boar taint (and
 CC improve meat quality) and for immunocastration
 XX Sequence 45 AA;
 SQ
 Query Match 86.8%; Score 211; DB 3; Length 45;
 Best Local Similarity 88.9%; Pred. No. 1.6e-22;
 Matches 40; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
 QY 1 TAKSKKPPSYTATYQFGSLSEIKGVIVHRLGEGVGHMSYGLRPG 45
 DB 1 TAKSKKPPSYTATYQFGSLSEIKGVIVHRLGEGVGHMSYGLRPG 45
 RESULT 9
 AAAY91172
 ID AAAY91172 standard; peptide; 45 AA.
 AC AAAY91172;
 DT 12-SEP-2003 (revised)
 DT 22-MAY-2000 (first entry)
 XX
 DE Modified WVF Th epitope/LHRH antigenic peptide, SEQ ID NO:52.
 XX
 KW Promiscuous T-cell epitope; measles virus F protein; WVF;
 KW hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;
 KW luteinizing hormone releasing hormone; LHRH; contraceptive; anticancer;
 KW somatostatin; growth promotion; CD4 receptor; HIV-1; antiviral; FMDV;
 KW foot and mouth disease virus; immunoglobulin E; IGE; anti-allergic;
 KW Plasmodium falciparum; circumsporozoite; antimalarial; CEMP;
 KW cholesterol ester transport protein; anti-arteriosclerotic.
 XX
 OS Measles virus.
 OS Ratus sp.
 OS Chimeric.
 OS
 PN MO9966957-A2.
 PD 29-DEC-1999.
 PF 21-JUN-1999; 99MO-US013975.
 PR 20-JUN-1998; 98US-00100412.
 XX

PA (UNBI-) UNITED BIOMEDICAL INC.
 XX Wang CY;
 PI WPI, 2000-160564/14.
 DR WPI, 2000-160564/14.
 XX
 PT New artificial T helper cell epitope and derived immunogens with target
 PT antigenic site, for immunization against e.g. malaria, arteriosclerosis
 PS or human immune deficiency virus.
 XX
 PS Example 1; Page 83; 129pp; English.
 XX
 CC The invention relates to novel promiscuous T helper cell epitopes (Th),
 CC and immunogenic peptides comprising the Th epitopes of the invention
 CC along with B cell epitopes. The Th epitopes and peptide immunogens
 CC containing them, are used to induce a T helper cell response,
 CC specifically against Plasmodium falciparum, cholesterol ester transport
 CC protein (CEP) or HIV epitopes, but more generally against any pathogen,
 CC immunoreactive self-antigen or tumour antigen. The Th epitopes and
 CC peptide immunogens may be used for prevention and/or treatment of
 CC infections (HIV, foot-and-mouth disease or malaria); for cancer
 CC immunotherapy; for inhibition of the action of luteinizing hormone
 CC releasing hormone (LHRH) for contraception, treatment of hormone-
 CC dependent cancer, prevention of boar taint in meat, and immunocastration
 CC ; for promoting the growth of animals; or for treating allergies or
 CC arteriosclerosis. Incorporation of a promiscuous Th (functional in
 CC induce a strong T helper cell-mediated immune response, resulting in
 CC production of antibodies against a target antigen. Th can replace carrier
 CC proteins and pathogen-derived T helper epitopes. Sequence AAAY91121
 CC represents a promiscuous T helper epitope from the measles virus F (WVF)
 CC protein and sequences AAAY91122-Y91142, AAAY9126 and AAAY91245-Y91246
 CC represent synthetic Th epitopes based on the WVF Th epitope. Sequence
 CC AAAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV)
 CC derived from this HBV epitope. AAAY91154-Y91155 are synthetic epitopes
 CC Y91244 are antigenic peptides comprising an LHRH sequence joined to a
 CC promiscuous Th epitope. AAAY91197 is the LHRH target antigenic peptide
 CC used in these LHRH antigenic peptides. AAAY91200 is somatostatin, and
 CC AAAY91201-Y91207 are antigenic peptides comprising somatostatin, and
 CC epitope. Somatostatin immunogens may be used to promote growth in
 CC livestock. AAAY91208 is a human CD4/CDR2-like domain antigenic site, and
 CC AAAY91209-Y90211 are MVA Th epitope/CD4/CDR2 antigenic peptides which may
 CC be used to prevent HIV infection of T cells. AAAY90212 is a modified
 CC version of a human IGE (immunoglobulin E) CH3 domain, and AAAY90213-Y90219
 CC are Th epitope/IGE CH3 antigenic peptides which may be used in the
 CC treatment of allergies. AAAY91220 is a peptide derived from foot and mouth
 CC disease virus (FMDV) VP1 capsid protein and AAAY91221-Y91222 comprise this
 CC circumsporozoite (CS) target antigen, and AAAY91223 comprise the CS
 CC antigen and an MVA Th epitope and may be used in a malaria vaccine.
 CC AAAY91228-Y91231 represent CEMP-derived peptides and AAAY91232-Y91241 are
 CC immunogens comprising a CEMP peptide and a Th epitope which may be used
 CC to prevent or treat arteriosclerosis and cardiovascular disease. AAAY91247
 CC and AAAY91252-Y91257 are HIV-1 neutralising B-cell epitopes, and AAAY91248-
 CC Y91251 and AAAY91258-Y91273 are antigenic peptides comprising MVA Th and
 CC HIV-1 B-cell epitope which may be used as a component in an anti-HIV-1
 CC vaccine. AAAY91198 and AAAY91199 are respectively an immunostimulatory
 CC both of which may optionally be used in the antigenic peptides of the
 CC invention. (Updated on 12-SEP-2003 to standardise OS field)
 XX
 SQ Sequence 45 AA;
 Query Match 86.8%; Score 211; DB 3; Length 45;
 Best Local Similarity 88.9%; Pred. No. 1.6e-22;
 Matches 40; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
 QY 1 TAKSKKPPSYTATYQFGSLSEIKGVIVHRLGEGVGHMSYGLRPG 45
 DB 1 TAKSKKPPSYTATYQFGSLSEIKGVIVHRLGEGVGHMSYGLRPG 45

RESULT 10
AAV68583
ID AAV68583 standard; peptide: 47 AA.
XX
AC AAV68583;
XX
DT 05-MAY-2000 (first entry)
XX
DE Peptide immunogen comprising a Th epitope and LHRH target antigen.
XX
KM Helper T cell epitope; F protein; Measles virus; peptide immunogen; LHRH;
KM luteinizing hormone-releasing hormone; spermatogenesis; ovulation;
KM oestrus; sexual development; sex hormone; promiscuous T helper epitope;
KM vaccine; contraceptive; hormone-dependent tumour; prostate cancer;
KM breast cancer; endometriosis; boar taint; meat quality; invasion domain;
KM immunocastration.
XX
OS Synthetic.
OS Yersinia sp.
OS Measles virus.
OS Undifferentiated.
XX
FH Key
FT Peptide
FT Peptide
FT Peptide
FT Peptide
FT Peptide
FT Peptide
XX
PN W09966952-A1.
XX
PD 29-DEC-1999.
XX
PF 21-JUN-1999; 99WC-US013960.
XX
PR 20-JUN-1998; 98US-00100414.
XX
PA (UNBI-) UNITED BIOMEDICAL INC.
XX
PI Wang CY;
XX
DR WPI; 2000-160562/14.
XX
PT New peptide immunogen containing luteinizing hormone-releasing hormone
PT antigen site and helper T cell epitope, for e.g. contraception and
PT treatment of cancer.
XX
PS Claim 9; Page 80; 102pp; English.
XX
CC The present sequence represents a peptide immunogen comprising an invasion
CC domain immunostimulatory peptide of Yersinia sp., a synthetic helper T
CC cell (Th) epitope and a target antigen, luteinizing hormone-releasing
CC hormone (LHRH). The synthetic Th epitope is derived from a structured
CC synthetic antigen library (SSAL) designated SSAL Th1. SSAL Th1 is
CC modelled after a promiscuous epitope taken from the F protein of the
CC Measles virus. The peptide immunogens cause induction of a specific
CC immune response to LHRH which is involved in regulation of a specific
CC spermatogenesis, ovulation, oestrus, sexual development and secretion of
CC sex hormones. Provision of a promiscuous T helper epitope (which is
CC functional in genetically diverse subjects) provides optimum
CC immunogenicity to the B cell epitopes of the target antigen and thus high
CC antibody titres against the target antigen. The peptide immunogens of the
CC invention are used to vaccinate against mammalian LHRH, for use as
CC (reversible) contraceptive; control of hormone-dependent tumours (cancer
CC of prostate or breast, also endometriosis); to prevent boar taint (and
CC improve meat quality) and for immunocastration
XX
SQ Sequence 47 AA;
Query Match 85.6%; Score 208; DB 3; Length 47;

Best Local Similarity 83.0%; Pred. No. 4,6e-22;
Matches 39; Conservative 4; Mismatches 2; Indels 2; Gaps 1;
Qy 1 TASSKPEPSYATYQFGSLSEIKGVIVHRLGV--GGEHWSYGLRPG 45
Db 1 TASSKPEPSYATYQFGSLSEIKGVIVHRLGVIVHRLGV--GGEHWSYGLRPG 47
RESULT 11
AAV91180
ID AAV91180 standard; peptide: 47 AA.
XX
AC AAV91180;
XX
DT 12-SEP-2003 (revised)
DT 22-MAY-2000 (first entry)
XX
DE Inv epitope/modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:60.
XX
KM Promiscuous T-cell epitope; measles virus F protein; MVF;
KM hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;
KM luteinizing hormone releasing hormone; LHRH; contraceptive; anticancer;
KM somatostatin; growth promotion; CD4 receptor; HIV-1; antiviral; PMDV;
KM foot and mouth disease virus; immunoglobulin E; IGE; anti-allergic;
KM Plasmodium falciparum; circumsporozoite; antimalarial; CERP;
KM cholesterol ester transport protein; anti-arteriosclerotic.
XX
OS Measles virus.
OS Rattus sp.
OS Yersinia sp.
OS Chimeric.
XX
PN W09966957-A2.
XX
PD 29-DEC-1999.
XX
PF 21-JUN-1999; 99WC-US013975.
XX
PR 20-JUN-1998; 98US-00100412.
XX
PA (UNBI-) UNITED BIOMEDICAL INC.
XX
PI Wang CY;
XX
DR WPI; 2000-160564/14.
XX
PT New artificial T helper cell epitope and derived immunogens with target
PT antigenic site, for immunization against e.g. malaria, arteriosclerosis
PT or human immune deficiency virus.
XX
PS Example 1; Page 86; 129pp; English.
XX
CC The invention relates to novel promiscuous T helper cell epitopes (Th),
CC and immunogenic peptides comprising the Th epitopes of the invention
CC along with B cell epitopes. The Th epitopes and peptide immunogens
CC containing them, are used to induce a T helper cell response,
CC specifically against Plasmodium falciparum, cholesterol ester transport
CC protein (CERP) or HIV epitopes, but more generally against any pathogen,
CC immunoreactive self-antigen or tumour antigen. The Th epitopes and
CC peptide immunogens may be used for prevention and/or treatment of
CC infections (HIV, foot-and-mouth disease or malaria); for cancer
CC immunotherapy; for inhibition of the action of luteinizing hormone
CC releasing hormone (LHRH) for contraception, treatment of hormone-
CC dependent cancer, prevention of boar taint in meat, and immunocastration
CC ; for promoting the growth of animals; or for treating allergies or
CC arteriosclerosis. Incorporation of a promiscuous Th (functional in
CC genetically diverse subjects) into an immunogen improves capacity to
CC induce a strong T helper cell-mediated immune response, resulting in
CC production of antibodies against a target antigen. Th can replace carrier
CC proteins and pathogen-derived T helper epitopes. Sequence AAV91121
CC represents a promiscuous T helper epitope from the measles virus F (MVF)
CC protein and sequences AAV91122-Y91142, AAV9126 and AAV91245-Y91246
CC represent synthetic Th epitopes based on the MVF Th epitope. Sequence

ID	AAV68586 standard; peptide; 47 AA.
XX	AAV68586;
AC	AAV68586;
DT	05-MAY-2000 (first entry)
XX	Peptide immunogen comprising a Th epitope and LHRH target antigen.
XX	Helper T cell epitope; F protein; Measles virus; peptide immunogen; LHRH;
KW	luteinizing hormone-releasing hormone; spermatogenesis; ovulation;
KW	oestrus; sexual development; sex hormone; promiscuous T helper epitope;
KW	vaccini; contraceptive; hormone-dependent tumour; prostate cancer;
KW	breast cancer; endometriosis; boar taint; meat quality; invasion domain;
KW	immunocastration.
XX	Synthetic.
OS	Yersinia sp.
OS	Measles virus.
OS	Unidentified.
XX	
FT	Key
FT	Location/Qualifiers
FT	Peptide
FT	1..16
FT	/note= "invasin domain AAV68565"
FT	17..35
FT	/note= "helper Th epitope AAV68553"
FT	36..37
FT	/note= "spacer"
FT	38..47
FT	/note= "LHRH antigenic epitope AAV68566"
XX	
XX	MO9966952-A1.
XX	
PD	29-DEC-1999.
XX	
PF	21-JUN-1999; 99MO-US013960.
XX	
PR	20-JUN-1998; 98US-00100414.
PA	(UNBI-) UNITED BIOMEDICAL INC.
XX	
P1	Wang CY;
DR	WPI; 2000-160562/14.
XX	
XX	New peptide immunogen containing luteinizing hormone-releasing hormone
PT	antigen site and helper T cell epitope, for e.g. contraception and
PT	treatment of cancer.
PS	Claim 9; Page 81; 102pp; English.
XX	
CC	The present sequence represents a peptide immunogen comprising an invasin
CC	domain immunostimulatory peptide of Yersinia sp., a synthetic helper T
CC	cell (Th) epitope and a target antigen, luteinizing hormone-releasing
CC	hormone (LHRH). The synthetic Th epitope is derived from a structured
CC	synthetic antigen library (SSAL) designated SSAL Th1. SSAL Th1 is
CC	modelled after a promiscuous epitope taken from the F protein of the
CC	Measles virus. The peptide immunogens cause induction of a specific
CC	immune response to LHRH which is involved in regulation of
CC	spermatogenesis, ovulation, oestrus, sexual development and secretion of
CC	sex hormones. Provision of a promiscuous T helper epitope (which is
CC	functional in genetically diverse subjects) provides optimum
CC	immunogenicity to the B cell epitopes of the target antigen and thus high
CC	antibody titres against the target antigen. The peptide immunogens of the
CC	invention are used to vaccinate against mammalian LHRH, for use as
CC	(reversible) contraceptive; control of hormone-dependent tumours (cancer
CC	of prostate or breast, also endometriosis); to prevent boar taint (and
CC	improve meat quality) and for immunocastration
XX	
XX	Sequence 47 AA;
XX	
XX	Query Match
XX	Best Local Similarity 80.9%; Score 203; DB 3; Length 47;
XX	Matches 38; Conservative 5; Mismatches 2; Indels 2; Gaps 1

Oy	TAKSKPEPSYATYQFGSEIKGVYVHRLGEV--GGHEWMSGLRAP 45
	:: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: ::
Dd	1 TAKSKFPSTATTATPQISMSKGVIVHMGMLFGGHEMSGLRAP 47
.	..
RESULT 14	
ID	AAY91183
XX	AAY91183 standard; peptide; 47 AA.
AC	AAY91183;
DT	12-SEP-2003 (revised)
D7	22-MAY-2000 (first entry)
XX	
De	Inw epitope/modified MVF Th epitope/LHRH antigenic peptide, SEQ ID NO:63
KM	Promiscuous T-cell epitope; measles virus F protein; MVF;
KW	hepatitis B virus surface antigen; HBV; immunogenic; B-cell epitope;
KW	luteinizing hormone releasing hormone; LHRH; contraceptive; anticancer;
KW	somatostatin; growth promotion; CD4 receptor; HIV-1; antiviral; FMDV;
KM	foot and mouth disease virus; immunoglobulin E; IgE; anti-allergic;
KM	Plasmodium falciparum; circumsporozoite; antimalarial; CEST;
KX	cholesterol ester transport protein; anti-arteriosclerotic.
OS	
OS	Measles virus.
OS	Rattus sp.
OS	Yersinia sp.
OS	Chimeric.
XX	
PN	MO9966957-A2.
PD	
PD	29-DEC-1999.
PF	21-JUN-1999; 99MO-US013975.
XX	
PR	20-JUN-1998; 98US-00100412.
XX	
PA	(UNBI-) UNITED BIOMEDICAL INC.
PI	
PI	Mang CY;
XX	
DR	WPJ; 2000-160564/14.
PT	
PT	New artificial T helper cell epitope and derived immunogens with target
PT	antigenic site, for immunization against e.g. malaria, arteriosclerosis
PT	or human immune deficiency virus.
XX	
PS	
PS	Example 1; Page 87; 129pp; English.
XX	
CC	The invention relates to novel promiscuous T helper cell epitopes (Th),
CC	and immunogenic peptides comprising the Th epitopes of the invention
CC	along with B cell epitopes. The Th epitopes and peptide immunogens
CC	containing them, are used to induce a T helper cell response,
CC	specifically against Plasmodium falciparum, cholesterol ester transport
CC	protein (CEMP) or HIV epitopes, but more generally against any pathogen,
CC	immunoreactive self-antigen or tumour antigen. The Th epitopes and
CC	peptide immunogens may be used for prevention and/or treatment of
CC	infections (HIV, foot-and-mouth disease or malaria); for cancer
CC	immunotherapy; for inhibition of the action of luteinizing hormone
CC	releasing hormone (LHRH) for contraception, treatment of hormone-
CC	dependent cancer, prevention of boar taint in meat, and immunocastration)
CC	; for promoting the growth of animals; or for treating allergies or
CC	arteriosclerosis. Incorporation of a promiscuous Th (functional in
CC	genetically diverse subjects) into an immunogen improves capacity to
CC	induce a strong T helper cell-mediated immune response, resulting in
CC	production of antibodies against a target antigen. Th can replace carrier
CC	proteins and pathogen-derived T helper epitopes. Sequence AAY91121
CC	represents a promiscuous T helper epitope from the measles virus F (MVFP)
CC	protein and sequences AAY91122-Y91142, AAY91226 and AAY91245-Y91246
CC	represent synthetic Th epitopes based on the MVF Th epitope. Sequence
CC	AAY91143 represents a promiscuous Th epitope from hepatitis B virus (HBV)
CC	surface antigen, and sequences AAY91144-Y91155 are synthetic epitopes

CC derived from this HBV epitope. AAY91156-Y91196, AAY91227 and AAY91242-
 CC Y91244 are antigenic peptides comprising an LHRH sequence joined to a
 CC promiscuous Th epitope. AAY91197 is the LHRH target antigenic peptide
 CC used in these LHRH antigenic peptides. AAY91200 is somatostatin, and
 CC AAY91201-Y91207 are antigenic peptides comprising somatostatin and a Th
 CC epitope. Somatostatin immunogens may be used to promote growth in
 CC livestock. AAY91208 is a human CD4 CDR2-like domain antigenic site, and
 CC AAY91209-Y90211 are MVA Th epitope/CD4 CDR2 antigenic peptides which may
 CC be used to prevent HIV infection of T cells. AAY90212 is a modified
 CC version of a human IGE (immunoglobulin E) CH3 domain, and AAY90213-Y90219
 CC are Th epitope/IGE CH3 antigenic peptides which may be used in the
 CC treatment of allergies. AAY91220 is a peptide derived from foot and mouth
 CC disease virus (FMDV) VP1 capsid protein and AAY91221-Y91222 comprise this
 CC peptide and a Th epitope. AAY91223 is a Plasmodium falciparum
 CC circumsporozoite (CS) target antigen, and AAY91224-Y91225 comprise the CS
 CC antigen and an MVA Th epitope and may be used in a malaria vaccine.
 CC AAY91228-Y91231 represent CEMP-derived peptides and AAY91232-Y91241 are
 CC immunogens comprising a CEMP peptide and a Th epitope which may be used
 CC to prevent or treat arteriosclerosis and cardiovascular disease. AAY91247
 CC and AAY91252-Y91257 are HIV-1 neutralizing B-cell epitopes, and AAY91248-
 CC Y91251 and AAY91258-Y91273 are antigenic peptides comprising MVA Th and
 CC HIV-1 B-cell epitope which may be used as a component in an anti-HIV-1
 CC vaccine. AAY91198 and AAY91199 are respectively an immunostimulatory
 CC invasion protein epitope from Yersinia species, and hinge spacer peptide,
 CC both of which may optionally be used in the antigenic peptides of the
 CC invention. (Updated on 12-SEP-2003 to standardise OS field)

XX
 SQ Sequence 47 AA;

Query Match 83.5%; Score 203; DB 3; Length 47;
 Best Local Similarity 80.9%; Pred. No. 2.4e-21;
 Matches 38; Conservative 5; Mismatches 2; Indels 2; Gaps 1;

QY 1 TAAKSKPPSYATATQFGGLSEIKGVYHRLGCV--GGEHMSYGLRPG 45
 1 TAAKSKPPSYATATQFGGLSEIKGVYHRLGCV--GGEHMSYGLRPG 47

RESULT 15
 AAY68571
 ID AAY68571 standard; peptide; 45 AA.

XX AC AAY68571;

XX DT 05-MAY-2000 (first entry)

XX DE Peptide immunogen comprising a Th epitope and LHRH target antigen.

XX KM Helper T cell epitope; F protein; Measles virus; peptide immunogen; LHRH;
 KM luteinizing hormone-releasing hormone; spermatogenesis; ovulation;
 KM oestrus; sexual development; sex hormone; promiscuous T helper epitope;
 KM vaccine; contraceptive; hormone-dependent tumour; prostate cancer;
 KM breast cancer; endometriosis; boar taint; meat quality; invasion domain;
 KM immunocastration.

XX OS Synthetic.
 OS Yersinia sp.
 OS Measles virus.
 OS Unidentified.

XX FH Key Location/Qualifiers

FT Peptide 1..16 /note= "invasion domain AAY68565"

FT Peptide 17..18 /note= "spacer"

FT Peptide 19..33 /note= "helper Th epitope AAY68542"

FT MISC-difference 20 /label= Ser, Thr

FT MISC-difference 23 /label= Lys, Arg

FT MISC-difference 24 /label= Gly, Thr

FT MISC-difference 28 /label= His, Thr
 FT MISC-difference 29 /label= Lys, Arg
 FT MISC-difference 32 /label= Gly, Thr
 FT Peptide 34..35 /note= "spacer"
 FT Peptide 36..45 /note= "LHRH antigenic epitope AAY68566"

XX PN WO966952-A1.

XX PD 29-DEC-1999.

XX PF 21-JUN-1999; 99WO-US013960.

XX PR 20-JUN-1998; 98US-00100414.

XX PA (UNB1-) UNITED BIOMEDICAL INC.

XX PI Wang CY;

XX PS WPI; 2000-160562/14.

PT New peptide immunogen containing luteinizing hormone-releasing hormone
 PT antigen site and helper T cell epitope, for e.g. contraception and
 PT treatment of cancer.

XX PS Claim 9; Page 69; 102pp; English.

XX CC The present sequence represents a peptide immunogen comprising an invasion
 CC domain immunostimulatory peptide of Yersinia sp., a synthetic helper T
 CC cell (Th) epitope and a target antigen, luteinizing hormone-releasing
 CC hormone (LHRH). The synthetic Th epitope is derived from a structured
 CC synthetic antigen library (SSAL) designated SSAL Th1. SSAL Th1 is
 CC modelled after a promiscuous epitope taken from the F protein of the
 CC Measles virus. The peptide immunogens cause induction of a specific
 CC immune response to LHRH which is involved in regulation of
 CC spermatogenesis, ovulation, oestrus, sexual development and secretion of
 CC sex hormones. Provision of a promiscuous T helper epitope (which is
 CC functional in genetically diverse subjects) provides optimum
 CC immunogenicity to the B cell epitopes of the target antigen and thus high
 CC antibody titres against the target antigen. The peptide immunogens of the
 CC invention are used to vaccinate against mammalian LHRH, for use as
 CC (reversible) contraceptive; control of hormone-dependent tumours (cancer
 CC of prostate or breast, also endometriosis); to prevent boar taint (and
 CC improve meat quality) and for immunocastration

SQ Sequence 45 AA;

Query Match 81.9%; Score 199; DB 3; Length 45;

Best Local Similarity 80.0%; Pred. No. 8.5e-21;
 Matches 36; Conservative 3; Mismatches 6; Indels 0; Gaps 0;

QY 1 TAAKSKPPSYATATQFGGLSEIKGVYHRLGCVGGEHMSYGLRPG 45
 1 TAAKSKPPSYATATQFGGLSEIKGVYHRLGCVGGEHMSYGLRPG 45

Search completed: February 8, 2005, 17:27:51
 Job time : 170 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: February 8, 2005, 17:25:08 ; Search time 43 Seconds
(without alignments)
78.121 Million cell updates/sec

Title: US-10-076-674A-9

Perfect score: 243
Sequence: 1 TASKSKFPSTATYFGSLGSL.....IVHREGVGGEHWSYGLRPG 45

Scoring table: BLOSUM62
Gapop 10.0, Gapext 0.5

Searched: 513545 seqs, 74649064 residues

Total number of hits satisfying chosen parameters: 513545

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: Issued Patents AA:*
1: /cgn2_6/ptodata/1/1aa/5A.COMB.pep:*
2: /cgn2_6/ptodata/1/1aa/5B.COMB.pep:*
3: /cgn2_6/ptodata/1/1aa/6A.COMB.pep:*
4: /cgn2_6/ptodata/1/1aa/6B.COMB.pep:*
5: /cgn2_6/ptodata/1/1aa/PTUS.COMB.pep:*
6: /cgn2_6/ptodata/1/1aa/backfile1.pep:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	ID	Description
1	243	100.0	US-08-446-692-33	Sequence 33, App1
2	243	100.0	US-08-488-351A-33	Sequence 33, App1
3	235	96.7	US-09-100-414B-45	Sequence 45, App1
4	235	96.7	US-09-303-323-45	Sequence 45, App1
5	235	96.7	US-09-701-588C-45	Sequence 45, App1
6	235	96.7	US-09-701-588C-45	Sequence 45, App1
7	216.5	89.1	US-08-446-692-40	Sequence 40, App1
8	216.5	89.1	US-08-488-351A-40	Sequence 40, App1
9	213	87.7	US-09-100-414B-57	Sequence 57, App1
10	213	87.7	US-09-303-323-57	Sequence 57, App1
11	213	87.7	US-09-701-588C-57	Sequence 57, App1
12	213	87.7	US-09-701-588C-57	Sequence 57, App1
13	211	86.8	US-09-100-414B-52	Sequence 52, App1
14	211	86.8	US-09-303-323-52	Sequence 52, App1
15	211	86.8	US-09-701-588C-52	Sequence 52, App1
16	211	86.8	US-09-701-588C-52	Sequence 52, App1
17	208	85.6	US-09-100-414B-60	Sequence 60, App1
18	208	85.6	US-09-303-323-60	Sequence 60, App1
19	208	85.6	US-09-701-588C-60	Sequence 60, App1
20	208	85.6	US-09-701-588C-60	Sequence 60, App1
21	206	84.8	US-09-100-414B-46	Sequence 46, App1
22	206	84.8	US-09-303-323-46	Sequence 46, App1
23	206	84.8	US-09-701-588C-46	Sequence 46, App1
24	206	84.8	US-09-701-588C-46	Sequence 46, App1
25	203	83.5	US-09-100-414B-63	Sequence 63, App1
26	203	83.5	US-09-303-323-63	Sequence 63, App1
27	203	83.5	US-09-701-588C-63	Sequence 63, App1

28	203	83.5	US-09-701-588C-63	Sequence 63, App1
29	187	77.0	US-09-100-414B-58	Sequence 58, App1
30	187	77.0	US-09-100-414B-62	Sequence 62, App1
31	187	77.0	US-09-303-323-58	Sequence 58, App1
32	187	77.0	US-09-303-323-62	Sequence 62, App1
33	187	77.0	US-09-701-588C-58	Sequence 58, App1
34	187	77.0	US-09-701-588C-58	Sequence 58, App1
35	187	77.0	US-09-701-588C-58	Sequence 58, App1
36	187	77.0	US-09-701-588C-62	Sequence 62, App1
37	180	74.1	US-08-446-692-55	Sequence 55, App1
38	180	74.1	US-08-488-351A-55	Sequence 55, App1
39	169.5	69.8	US-08-446-692-37	Sequence 37, App1
40	169.5	69.8	US-08-488-351A-37	Sequence 37, App1
41	159	65.4	US-08-446-692-36	Sequence 36, App1
42	159	65.4	US-08-488-351A-36	Sequence 36, App1
43	156.5	64.4	US-08-446-692-34	Sequence 34, App1
44	156.5	64.4	US-08-488-351A-34	Sequence 34, App1
45	156	64.2	US-08-446-692-32	Sequence 32, App1

ALIGNMENTS

RESULT 1
US-08-446-692-33
Sequence 33, Application US/08446692
Patent No. 5759551
GENERAL INFORMATION:
APPLICANT: Ladd, Anna
APPLICANT: Wang, Chang YI
APPLICANT: Zamb, Timothy
TITLE OF INVENTION: Immunogenic LHRH peptide constructs
TITLE OF INVENTION: and synthetic universal immune stimulants for vaccines
NUMBER OF SEQUENCES: 114
CORRESPONDENCE ADDRESS:
ADDRESSEE: Maria C.H. Lin
STREET: 345 Park Avenue
CITY: New York
STATE: NY
COUNTRY: US
ZIP: 10154-0053
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/446,692
FILING DATE: 7-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Maria C.H. Lin
REGISTRATION NUMBER: 29,323
REFERENCE/DOCKET NUMBER: 1151-4146 US2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212)415-8745
TELEFAX: (516)751-6849
INFORMATION FOR SEQ ID NO: 33:
SEQUENCE CHARACTERISTICS:
LENGTH: 45 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-08-446-692-33

Query Match 100.0%; Score 243; DB 1; Length 45;
Best Local Similarity 100.0%; Pred. No. 2.3e-28; Indels 0;
Matches 45; Conservative 0; Mismatches 0; Gaps 0;

Qy 1 TASKSKFPSTATYFGSLGSLIVHREGVGGEHWSYGLRPG 45
Db 1 TASKSKFPSTATYFGSLGSLIVHREGVGGEHWSYGLRPG 45

RESULT 2

US-08-488-351A-33

; Sequence 33, Application US/08488351A
; Patent No. 5843446

GENERAL INFORMATION:

; APPLICANT: Ladd, Anna
; APPLICANT: Wang, Chang Yi
; TITLE OF INVENTION: Immunogenic LHRH peptide constructs
; NUMBER OF SEQUENCES: 114
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Maria C.H. Lin
; STREET: 345 Park Avenue
; CITY: New York
; STATE: NY
; COUNTRY: US
; ZIP: 10154-0053

COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/488,351A
; FILING DATE: 7-JUN-1995
; CLASSIFICATION: 424

PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/446,692
; FILING DATE: 7-JUN-1995
; CLASSIFICATION: 424
; PRIOR APPLICATION NUMBER: 424
; FILING DATE: 14-APR-1994
; CLASSIFICATION: 424

APPLICATION DATA:

; APPLICATION NUMBER: US 08/057,166
; FILING DATE: 27-APR-1992
; CLASSIFICATION: 424
; ATTORNEY/AGENT INFORMATION:
; NAME: Maria C.H. Lin

REGISTRATION NUMBER: 29,323

; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 751-6849

TELEFAX: (516) 751-6849

INFORMATION FOR SEQ ID NO: 33:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 45 amino acids
; TYPE: amino acid
; TOPOLOGY: linear; MOLECULE TYPE: peptide
US-08-488-351A-33

Query Match 100.0%; Score 243; DB 2; Length 45;

Best Local Similarity 100.0%; Pred. No. 2.3e-28;

Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TAKSKKPPSYATYQFGSLSEIKGVIVHRLGVGGEHMSYGLRPG 45

Db 1 TAKSKKPPSYATYQFGSLSEIKGVIVHRLGVGGEHMSYGLRPG 45

RESULT 3

US-09-100-414B-45

; Sequence 45, Application US/09100414B
; Patent No. 6025468

GENERAL INFORMATION:

; APPLICANT: Wang, Chang Yi
; TITLE OF INVENTION: NOVEL LHRH PEPTIDE
; NUMBER OF SEQUENCES: 106

CORRESPONDENCE ADDRESS:

; ADDRESSEE: Morgan & Finnegan, L.L.P.
; STREET: 345 Park Avenue
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10154-0054

COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC Windows
; SOFTWARE: Word 97
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/100,414B
; FILING DATE: 20-JUNE-1998
; CLASSIFICATION: 424

ATTORNEY/AGENT INFORMATION:

; NAME: Maria H. Lin
; REGISTRATION NUMBER: 29,323
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 212-758-4800
; TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 45:

; SEQUENCE CHARACTERISTICS:
; LENGTH: 45 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-09-100-414B-45

Query Match

Best Local Similarity 96.7%; Score 235; DB 3; Length 45;

Matches 41; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TAKSKKPPSYATYQFGSLSEIKGVIVHRLGVGGEHMSYGLRPG 45

Db 1 TAKSKKPPSYATYQFGSLSEIKGVIVHRLGVGGEHMSYGLRPG 45

RESULT 4

US-09-303-323-45

; Sequence 45, Application US/09303323
; Patent No. 6228987

GENERAL INFORMATION:

; APPLICANT: Wang, Chang Yi
; TITLE OF INVENTION: NOVEL LHRH PEPTIDE
; NUMBER OF SEQUENCES: 106
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Morgan & Finnegan, L.L.P.
; STREET: 345 Park Avenue
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10154-0054

COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC Windows
; SOFTWARE: Word 97
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/303,323
; FILING DATE: 30-APR-1999
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 09/100,414
; FILING DATE: 20-JUNE-1998

ATTORNEY/AGENT INFORMATION:

; NAME: Maria H. Lin
; REGISTRATION NUMBER: 29,323
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 212-758-4800
; TELEFAX: 212-751-6849

TELEPHONE: 212-758-4800
TELEFAX: 212-751-6849
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 45 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-303-323-45

Query Match 96.7%; Score 235; DB 3; Length 45;
Best Local Similarity 91.1%; Pred. No. 3.4e-27;
Matches 41; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAAKKEPSTATYATQFGSLSEIKGVYHRLGEGVGEHWSYGLRPG 45
DB 1 TAAKKEPSTATYATQFGSLSEIKGVYHRLGEGVGEHWSYGLRPG 45

RESULT 5

US-09-770-014-45
Sequence 45, Application US/09770014
Patent No. 6559282
GENERAL INFORMATION:
APPLICANT: Wang, Chang YI
TITLE OF INVENTION: NOVEL LHRH PEPTIDE
TITLE OF INVENTION: IMMUNOGENS
NUMBER OF SEQUENCES: 106
CORRESPONDENCE ADDRESS:
ADDRESSEE: Morgan & Finnegan, L.L.P.
STREET: 345 Park Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10154-0054
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC Windows
SOFTWARE: Word 97
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/770,014
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/100,414
FILING DATE: 20-JUNE-1998
ATTORNEY/AGENT INFORMATION:
NAME: Maria H. Lin
REGISTRATION NUMBER: 29,323
REFERENCE/DOCKET NUMBER: 1151-4157
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-758-4800
TELEFAX: 212-751-6849
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 45 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-770-014-45

Query Match 96.7%; Score 235; DB 4; Length 45;
Best Local Similarity 91.1%; Pred. No. 3.4e-27;
Matches 41; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAAKKEPSTATYATQFGSLSEIKGVYHRLGEGVGEHWSYGLRPG 45
DB 1 TAAKKEPSTATYATQFGSLSEIKGVYHRLGEGVGEHWSYGLRPG 45

RESULT 6
US-09-701-588C-45

Sequence 45, Application US/09701588C
Patent No. 6713301
GENERAL INFORMATION:
APPLICANT: UNITED BIOMEDICAL INC., ET AL.
TITLE OF INVENTION: ARTIFICIAL T HELPER CELL
EPITOPES AS IMMUNE STIMULATORS FOR SYNTHETIC
PEPTIDE IMMUNOGENS
NUMBER OF SEQUENCES: 151
CORRESPONDENCE ADDRESS:
ADDRESSEE: Morgan & Finnegan, L.L.P.
STREET: 345 Park Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10154-0054
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC Windows
SOFTWARE: Word 97
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/701,588C
FILING DATE: 29-No. 6713301-2002
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 09/100,414
FILING DATE: 20-JUNE-1998
ATTORNEY/AGENT INFORMATION:
NAME: Maria H. Lin
REGISTRATION NUMBER: 29,323
REFERENCE/DOCKET NUMBER: 1151-4158PC1
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-758-4800
TELEFAX: 212-751-6849
INFORMATION FOR SEQ ID NO: 45:
SEQUENCE CHARACTERISTICS:
LENGTH: 45 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 45:
US-09-701-588C-45

Query Match 96.7%; Score 235; DB 4; Length 45;
Best Local Similarity 91.1%; Pred. No. 3.4e-27;
Matches 41; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAAKKEPSTATYATQFGSLSEIKGVYHRLGEGVGEHWSYGLRPG 45
DB 1 TAAKKEPSTATYATQFGSLSEIKGVYHRLGEGVGEHWSYGLRPG 45

RESULT 7

US-08-446-692-40
Sequence 40, Application US/08446692
Patent No. 5759551
GENERAL INFORMATION:
APPLICANT: Ladd, Anna
APPLICANT: Wang, Chang YI
APPLICANT: Zamb, Timothy
TITLE OF INVENTION: Immunogenic LHRH peptide constructs
TITLE OF INVENTION: and synthetic universal immune stimulators for vaccines
NUMBER OF SEQUENCES: 114
CORRESPONDENCE ADDRESS:
ADDRESSEE: Maria C.H. Lin
STREET: 345 Park Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10154-0053
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/446, 692
FILING DATE: 7-JUN-1995
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Maria C.H. Lin
REGISTRATION NUMBER: 29,323
REFERENCE/DOCKET NUMBER: 1151-4146 US2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 415-8745
TELEFAX: (516) 751-6849
INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 46 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 19
OTHER INFORMATION: /note="D0.50;R0.50"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 20
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 22
OTHER INFORMATION: /note="E0.50;D0.50"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 23
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 24
OTHER INFORMATION: /note="K0.50;R0.50"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 26
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 27
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 28
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
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NAME/KEY: Modified-site
LOCATION: 30
OTHER INFORMATION: /note="X0.50;R0.50"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 31
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 32
OTHER INFORMATION: /note="E0.50;D0.50"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 34
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
US-08-446-692-40

Query Match 89.1%; Score 216.5; DB 1; Length 46;
Best Local Similarity 84.8%; Pred. No. 1.8e-24;
Matches 39; Conservative 6; Mismatches 0; Indels 1; Gaps 1;

Db 1 TATSKRPPSYTATYQFGG-LSEIKGVIYHRLGVGGEHMSYGLRPG 45
1 TATSKRPPSYTATYQFGGDLSELKGLLHKLKLEGLGGEHMSYGLRPG 46
RESULT 8
US-08-488-351A-40
Sequence 40, Application US/08488351A
Patent No. 5843446
GENERAL INFORMATION:
APPLICANT: Ladd, Anna
APPLICANT: Wang, Chang Yi
APPLICANT: Zamb, Timothy
TITLE OF INVENTION: Immunogenic linear peptide constructs
TITLE OF INVENTION: and synthetic universal immune stimulators for vaccines
NUMBER OF SEQUENCES: 114
CORRESPONDENCE ADDRESS:
ADDRESSEE: Maria C.H. Lin
STREET: 345 Park Avenue
CITY: New York
STATE: NY
COUNTRY: US
ZIP: 10154-0053
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
FILING DATE: 7-JUN-1995
APPLICATION NUMBER: US/08/488,351A
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/446,692
FILING DATE: 7-JUN-1995
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/229,275
FILING DATE: 14-APR-1994
CLASSIFICATION: 424
APPLICATION DATA:
APPLICATION NUMBER: US 08/057,166
FILING DATE: 27-APR-1992
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Maria C.H. Lin
REGISTRATION NUMBER: 29,323
REFERENCE/DOCKET NUMBER: 1151-4146 US2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 415-8745
TELEFAX: (516) 751-6849
INFORMATION FOR SEQ ID NO: 40:
SEQUENCE CHARACTERISTICS:
LENGTH: 46 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
FEATURE:
NAME/KEY: Modified-site
LOCATION: 19
OTHER INFORMATION: /note="D0.50;E0.50"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 20
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 22
OTHER INFORMATION: /note="E0.50;D0.50"
FEATURE:
NAME/KEY: Modified-site
LOCATION: 23
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"

FEATURE: NAME/KEY: Modified-site
LOCATION: 24
OTHER INFORMATION: /note="K0.50;R0.50"
FEATURE: NAME/KEY: Modified-site
LOCATION: 26
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE: NAME/KEY: Modified-site
LOCATION: 27
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE: NAME/KEY: Modified-site
LOCATION: 28
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE: NAME/KEY: Modified-site
LOCATION: 30
OTHER INFORMATION: /note="K0.50;R0.50"
FEATURE: NAME/KEY: Modified-site
LOCATION: 31
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
FEATURE: NAME/KEY: Modified-site
LOCATION: 32
OTHER INFORMATION: /note="E0.50;D0.50"
FEATURE: NAME/KEY: Modified-site
LOCATION: 34
OTHER INFORMATION: /note="L0.25;I0.25;V0.25;F0.25"
US-08-488-351A-40

Query Match 89.1%; Score 216.5; DB 2; Length 46;
Best Local Similarity 84.8%; Pred. No. 1.8e-24;
Matches 39; Conservative 6; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TAKSKKPPSYTATYQFGG--LSEIKGYIYHRLGCV--GGHWSYGLRPG 45
Db 1 TAKSKKPPSYTATYQFGGDLSELKGLLHLRLGGLGSHWSYGLRPG 46

RESULT 9
US-09-100-414B-57
Sequence 57, Application US/09100414B
Patent No. 6025468
GENERAL INFORMATION:
APPLICANT: Wang, Chang Y1
TITLE OF INVENTION: NOVEL LHRH PEPTIDE
TITLE OF INVENTION: IMMUNOGENS
NUMBER OF SEQUENCES: 106
CORRESPONDENCE ADDRESS:
ADDRESSEE: Morgan & Flinnegan, L.L.P.
STREET: 345 Park Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10154-0054
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC Windows
SOFTWARE: Word 97
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/100,414B
FILING DATE: 20-JUNE-1998
CLASSIFICATION: 424
ATTORNEY/AGENT INFORMATION:
NAME: Maria H. Lin
REGISTRATION NUMBER: 29,323
REFERENCE/DOCKET NUMBER: 1151-4157
TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800
TELEFAX: 212-751-6849
INFORMATION FOR SEQ ID NO: 57:
SEQUENCE CHARACTERISTICS:
LENGTH: 49 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-100-414B-57

Query Match 87.7%; Score 213; DB 3; Length 49;
Best Local Similarity 83.7%; Pred. No. 6.3e-24;
Matches 41; Conservative 4; Mismatches 0; Indels 4; Gaps 2;

Qy 1 TAKSKKPPSYTATYQFGG--LSEIKGYIYHRLGCV--GGHWSYGLRPG 45
Db 1 TAKSKKPPSYTATYQFGGISISIKGYIYHRLGGLFGSHWSYGLRPG 49

RESULT 10
US-09-303-323-57
Sequence 57, Application US/09303323
Patent No. 6228987
GENERAL INFORMATION:
APPLICANT: Wang, Chang Y1
TITLE OF INVENTION: NOVEL LHRH PEPTIDE
TITLE OF INVENTION: IMMUNOGENS
NUMBER OF SEQUENCES: 106
CORRESPONDENCE ADDRESS:
ADDRESSEE: Morgan & Flinnegan, L.L.P.
STREET: 345 Park Avenue
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10154-0054
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC Windows
SOFTWARE: Word 97
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/303,323
FILING DATE: 30-APR-1998
CLASSIFICATION:
PRIOR APPLICATION:
APPLICATION NUMBER: 09/100,414
FILING DATE: 20-JUNE-1998
ATTORNEY/AGENT INFORMATION:
NAME: Maria H. Lin
REGISTRATION NUMBER: 29,323
REFERENCE/DOCKET NUMBER: 1151-4157
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-758-4800
TELEFAX: 212-751-6849
INFORMATION FOR SEQ ID NO: 57:
SEQUENCE CHARACTERISTICS:
LENGTH: 49 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-303-323-57

Query Match 87.7%; Score 213; DB 3; Length 49;
Best Local Similarity 83.7%; Pred. No. 6.3e-24;
Matches 41; Conservative 4; Mismatches 0; Indels 4; Gaps 2;

Qy 1 TAKSKKPPSYTATYQFGG--LSEIKGYIYHRLGCV--GGHWSYGLRPG 45
Db 1 TAKSKKPPSYTATYQFGGISISIKGYIYHRLGGLFGSHWSYGLRPG 49

RESULT 11
US-09-770-014-57

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/ Sequence 57, Application US/09770014
/ Patent No. 6559282
/ GENERAL INFORMATION:
/ APPLICANT: Wang, Chang Yi
/ TITLE OF INVENTION: NOVEL LHRH PEPTIDE
/ NUMBER OF INVENTION: IMMUNOGENS
/ CORRESPONDENCE ADDRESSES:
/ ADDRESSEE: Morgan & Finnegan, L.L.P.
/ STREET: 345 Park Avenue
/ CITY: New York
/ STATE: NY
/ COUNTRY: USA
/ ZIP: 10154-0054
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Floppy disk
/ COMPUTER: IBM PC compatible
/ OPERATING SYSTEM: PC Windows
/ SOFTWARE: Word 97
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/09/770,014
/ FILING DATE:
/ CLASSIFICATION:
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: 09/100,414
/ FILING DATE: 20-JUNE-1998
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Maria H. Lin
/ REGISTRATION NUMBER: 29,323
/ REFERENCE/DOCKET NUMBER: 1151-4157
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 212-758-4800
/ TELEFAX: 212-751-6849
/ INFORMATION FOR SEQ ID NO: 57:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 49 amino acids
/ TYPE: amino acid
/ TOPOLOGY: linear
/ MOLECULE TYPE: peptide
/ US-09-770-014-57

Query Match      87.7%; Score 213; DB 4; Length 49;
Best Local Similarity 83.7%; Pred. No. 6,3e-24;
Matches 41; Conservative 4; Mismatches 0; Indels 4; Gaps 2;

OY 1 TAKSKKPSYATYQFGG--LSEIKGVIVHRLGV--GGEHWSYGLRPG 45
DB 1 TAKSKKPSYATYQFGGISSEIKGVIVHKLGLFGGEHWSYGLRPG 49

RESULT 12
US-09-701-588C-57
/ Sequence 57, Application US/09701588C
/ Patent No. 6713301
/ GENERAL INFORMATION:
/ APPLICANT: UNITED BIOMEDICAL INC., ET AL.
/ TITLE OF INVENTION: ARTIFICIAL T HELPER CELL
/ EPTIOPES AS IMMUNE STIMULATORS FOR SYNTHETIC
/ NUMBER OF SEQUENCES: 151
/ CORRESPONDENCE ADDRESSES:
/ ADDRESSEE: Morgan & Finnegan, L.L.P.
/ STREET: 345 Park Avenue
/ CITY: New York
/ STATE: NY
/ COUNTRY: USA
/ ZIP: 10154-0054
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Floppy disk
/ COMPUTER: IBM PC compatible
/ OPERATING SYSTEM: PC Windows
/ SOFTWARE: Word 97
/ CURRENT APPLICATION DATA:
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/ APPLICATION NUMBER: US/09/701,588C
/ FILING DATE: 29-NOV. 6713301-2002
/ CLASSIFICATION: <Unknown>
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: 09/100,414
/ FILING DATE: 20-JUNE-1998
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Maria H. Lin
/ REGISTRATION NUMBER: 29,323
/ REFERENCE/DOCKET NUMBER: 1151-4158PCL
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 212-758-4800
/ TELEFAX: 212-751-6849
/ INFORMATION FOR SEQ ID NO: 57:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 49 amino acids
/ TYPE: amino acid
/ TOPOLOGY: linear
/ MOLECULE TYPE: peptide
/ SEQUENCE DESCRIPTION: SEQ ID NO: 57:
/ US-09-701-588C-57

Query Match      87.7%; Score 213; DB 4; Length 49;
Best Local Similarity 83.7%; Pred. No. 6,3e-24;
Matches 41; Conservative 4; Mismatches 0; Indels 4; Gaps 2;

OY 1 TAKSKKPSYATYQFGG--LSEIKGVIVHRLGV--GGEHWSYGLRPG 45
DB 1 TAKSKKPSYATYQFGGISSEIKGVIVHKLGLFGGEHWSYGLRPG 49

RESULT 13
US-09-100-414B-52
/ Sequence 52, Application US/09100414B
/ Patent No. 6025468
/ GENERAL INFORMATION:
/ APPLICANT: Wang, Chang Yi
/ TITLE OF INVENTION: NOVEL LHRH PEPTIDE
/ NUMBER OF SEQUENCES: 106
/ CORRESPONDENCE ADDRESSES:
/ ADDRESSEE: Morgan & Finnegan, L.L.P.
/ STREET: 345 Park Avenue
/ CITY: New York
/ STATE: NY
/ COUNTRY: USA
/ ZIP: 10154-0054
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Floppy disk
/ COMPUTER: IBM PC compatible
/ OPERATING SYSTEM: PC Windows
/ SOFTWARE: Word 97
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/09/100,414B
/ FILING DATE: 20-JUNE-1998
/ CLASSIFICATION: 424
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Maria H. Lin
/ REGISTRATION NUMBER: 29,323
/ REFERENCE/DOCKET NUMBER: 1151-4157
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 212-758-4800
/ TELEFAX: 212-751-6849
/ INFORMATION FOR SEQ ID NO: 52:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 45 amino acids
/ TYPE: amino acid
/ TOPOLOGY: linear
/ MOLECULE TYPE: peptide
/ US-09-100-414B-52

Query Match      86.8%; Score 211; DB 3; Length 45;
Best Local Similarity 88.9%; Pred. No. 1.1e-23;
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Matches 40; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 TAKSKKPSTYATYQFGLSBKGVYHRLGVGGEHMSYGLRPG 45
Db 1 TAKSKKPSTYATYQFGLSBKGVYHRLGVGGEHMSYGLRPG 45

RESULT 14

US-09-303-323-52
Sequence 52, Application US/09303323

Patent No. 6228987

GENERAL INFORMATION:

APPLICANT: Wang, Chang Yi

TITLE OF INVENTION: NOVEL LHRH PEPTIDE

NUMBER OF SEQUENCES: 106

CORRESPONDENCE ADDRESS:

ADDRESSEE: Morgan & Finnegan, L.L.P.

STREET: 345 Park Avenue

CITY: New York

STATE: NY

COUNTRY: USA

ZIP: 10154-0054

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC Windows

SOFTWARE: Word 97

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/303,323

FILING DATE: 30-APR-1999

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 09/100,414

FILING DATE: 20-JUNE-1998

ATTORNEY/AGENT INFORMATION:

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 1151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 52:

SEQUENCE CHARACTERISTICS:

LENGTH: 45 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-303-323-52

Query Match 86.8%; Score 211; DB 3; Length 45;
Best Local Similarity 88.9%; Pred. No. 1,1e-23;
Matches 40; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 TAKSKKPSTYATYQFGLSBKGVYHRLGVGGEHMSYGLRPG 45
Db 1 TAKSKKPSTYATYQFGLSBKGVYHRLGVGGEHMSYGLRPG 45

RESULT 15

US-09-770-014-52
Sequence 52, Application US/09770014

Patent No. 6559282

GENERAL INFORMATION:

APPLICANT: Wang, Chang Yi

TITLE OF INVENTION: NOVEL LHRH PEPTIDE

NUMBER OF SEQUENCES: 106

CORRESPONDENCE ADDRESS:

ADDRESSEE: Morgan & Finnegan, L.L.P.

STREET: 345 Park Avenue

CITY: New York

STATE: NY

COUNTRY: USA

ZIP: 10154-0054

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC Windows

SOFTWARE: Word 97

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/770,014

FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 09/100,414

FILING DATE: 20-JUNE-1998

ATTORNEY/AGENT INFORMATION:

NAME: Maria H. Lin

REGISTRATION NUMBER: 29,323

REFERENCE/DOCKET NUMBER: 1151-4157

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-758-4800

TELEFAX: 212-751-6849

INFORMATION FOR SEQ ID NO: 52:

SEQUENCE CHARACTERISTICS:

LENGTH: 45 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: peptide

US-09-770-014-52

Query Match 86.8%; Score 211; DB 4; Length 45;
Best Local Similarity 88.9%; Pred. No. 1,1e-23;
Matches 40; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 TAKSKKPSTYATYQFGLSBKGVYHRLGVGGEHMSYGLRPG 45
Db 1 TAKSKKPSTYATYQFGLSBKGVYHRLGVGGEHMSYGLRPG 45

Search completed: February 8, 2005, 17:34:37
Job time: 44 secs

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OM protein - protein search, using sw model

Run on: February 8, 2005, 17:21:18 ; Search time 128 Seconds

(without alignments)

114.505 Million cell updates/sec

Title: US-10-076-674A-9

Sequence: 1 TAKSKKPPSYTATYQFGGLS.....IVHRLBVGGEHMSYGLRPG 45

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Total number of hits satisfying chosen parameters: 1373511

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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Database :

Published Applications AA:*

- 1: /cgn2_6/ptodata/2/pubppaa/US07_PUBCOMB.pep.*
- 2: /cgn2_6/ptodata/2/pubppaa/PCT_NEW_PUB.pep.*
- 3: /cgn2_6/ptodata/2/pubppaa/US06_NEW_PUB.pep.*
- 4: /cgn2_6/ptodata/2/pubppaa/US06_PUBCOMB.pep.*
- 5: /cgn2_6/ptodata/2/pubppaa/US07_NEW_PUB.pep.*
- 6: /cgn2_6/ptodata/2/pubppaa/PCTUS_PUBCOMB.pep.*
- 7: /cgn2_6/ptodata/2/pubppaa/US08_NEW_PUB.pep.*
- 8: /cgn2_6/ptodata/2/pubppaa/US08_PUBCOMB.pep.*
- 9: /cgn2_6/ptodata/2/pubppaa/US09_PUBCOMB.pep.*
- 10: /cgn2_6/ptodata/2/pubppaa/US09C_PUBCOMB.pep.*
- 11: /cgn2_6/ptodata/2/pubppaa/US09C_PUBCOMB.pep.*
- 12: /cgn2_6/ptodata/2/pubppaa/US09C_PUBCOMB.pep.*
- 13: /cgn2_6/ptodata/2/pubppaa/US10_PUBCOMB.pep.*
- 14: /cgn2_6/ptodata/2/pubppaa/US10C_PUBCOMB.pep.*
- 15: /cgn2_6/ptodata/2/pubppaa/US10C_PUBCOMB.pep.*
- 16: /cgn2_6/ptodata/2/pubppaa/US10C_PUBCOMB.pep.*
- 17: /cgn2_6/ptodata/2/pubppaa/US10C_PUBCOMB.pep.*
- 18: /cgn2_6/ptodata/2/pubppaa/US11_NEW_PUB.pep.*
- 19: /cgn2_6/ptodata/2/pubppaa/US11_NEW_PUB.pep.*
- 20: /cgn2_6/ptodata/2/pubppaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	243	100.0	45	14	US-10-076-674A-9
2	243	100.0	45	15	US-10-355-161A-9
3	157	64.6	45	15	US-10-076-674-8
4	157	64.6	45	15	US-10-355-161A-8
5	116	47.7	31	9	US-09-848-834A-9
6	116	47.7	47	9	US-09-848-834A-17
7	84	34.6	16	10	US-09-747-802-72
8	84	34.6	16	10	US-10-296-734-1467
9	84	34.6	16	16	US-10-789-619-72
10	84	34.6	25	15	US-10-411-544-32
11	79	32.5	36	14	US-10-351-641-505
12	79	32.5	438	15	US-10-267-682-105
13	79	32.5	438	15	US-10-267-748-105

14	79	32.5	550	9	US-09-873-233A-18	Sequence 18, App1
15	79	32.5	550	9	US-09-873-233A-20	Sequence 20, App1
16	72	29.6	15	10	US-09-747-802-16	Sequence 16, App1
17	72	29.6	15	10	US-09-747-802-30	Sequence 30, App1
18	72	29.6	15	10	US-09-865-294-8	Sequence 8, App1
19	72	29.6	15	10	US-09-865-294-22	Sequence 22, App1
20	72	29.6	15	14	US-10-261-446-20	Sequence 20, App1
21	72	29.6	15	15	US-10-411-544-10	Sequence 10, App1
22	72	29.6	15	15	US-10-261-446-20	Sequence 20, App1
23	72	29.6	15	16	US-10-789-619-16	Sequence 16, App1
24	72	29.6	15	16	US-10-789-619-30	Sequence 30, App1
25	72	29.6	15	16	US-10-782-234-20	Sequence 20, App1
26	72	29.6	16	9	US-09-848-834A-8	Sequence 8, App1
27	72	29.6	18	14	US-10-351-641-1148	Sequence 14, App1
28	72	29.6	34	9	US-09-848-834A-13	Sequence 13, App1
29	72	29.6	438	15	US-10-267-682-93	Sequence 93, App1
30	72	29.6	438	15	US-10-267-748-93	Sequence 93, App1
31	72	29.6	662	15	US-09-951-061A-141	Sequence 141, App1
32	72	29.6	662	15	US-10-670-695-36	Sequence 36, App1
33	71	29.2	284	15	US-10-358-083-6	Sequence 6, App1
34	71	29.2	600	15	US-10-282-122A-45023	Sequence 45023, App1
35	70	28.8	40	14	US-10-223-711-11	Sequence 11, App1
36	70	28.8	986	9	US-09-870-759-33	Sequence 33, App1
37	70	28.8	986	10	US-09-751-708A-33	Sequence 33, App1
38	69	28.4	15	10	US-09-747-802-37	Sequence 37, App1
39	69	28.4	15	10	US-09-865-294-29	Sequence 29, App1
40	69	28.4	15	16	US-10-789-619-37	Sequence 37, App1
41	69	28.4	19	10	US-09-747-802-48	Sequence 48, App1
42	69	28.4	19	10	US-09-865-294-40	Sequence 40, App1
43	69	28.4	19	16	US-10-789-619-48	Sequence 48, App1
44	69	28.4	20	10	US-09-964-201A-29	Sequence 29, App1
45	69	28.4	20	10	US-09-964-201A-31	Sequence 31, App1

ALIGNMENTS

RESULT 1
US-10-076-674-9
; Sequence 9, Application US/10076674
; Publication No. US20030165478A1
; GENERAL INFORMATION:
; APPLICANT: Sokoll, Kenneth K.
; TITLE OR INVENTION: Stabilized Synthetic Immunogen Delivery System
; FILE REFERENCE: Immunogen Delivery System
; CURRENT APPLICATION NUMBER: US/10/076,674
; CURRENT FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 9
; LENGTH: 45
; TYPE: PRT
; ORGANISM: Human
US-10-076-674-9

Query Match 100.0%; Score 243; DB 14; Length 45;
Best Local Similarly 100.0%; Pred. No. 3.3e-26;
Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPPSYTATYQFGGLSEIKGVIVHRLBVGGEHMSYGLRPG 45
DB 1 TAKSKKPPSYTATYQFGGLSEIKGVIVHRLBVGGEHMSYGLRPG 45

RESULT 2
US-10-355-161A-9
; Sequence 9, Application US/10355161A
; Publication No. US2004009897A1
; GENERAL INFORMATION:
; APPLICANT: Sokoll, Kenneth K.
; TITLE OR INVENTION: Stabilized Synthetic Immunogen Delivery System
; FILE REFERENCE: Immunogen Delivery System
; CURRENT APPLICATION NUMBER: US/10/355,161A

;; CURRENT FILING DATE: 2003-01-31
;; PRIOR APPLICATION NUMBER: US 10/076674
;; PRIOR FILING DATE: 2002-02-14
;; NUMBER OF SEQ ID NOS: 13
;; SOFTWARE: PatentIn version 3.1
;; SEQ ID NO 9
;; LENGTH: 45
;; TYPE: PRT
;; ORGANISM: Human
US-10-355-161A-9

Query Match
Best Local Similarity 100.0%; Score 243; DB 15; Length 45;
Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAAKSKFPSTATYQFGGLSEIKGIVYHRLGVGGHMSYGLRPG 45
DB 1 TAAKSKFPSTATYQFGGLSEIKGIVYHRLGVGGHMSYGLRPG 45

RESULT 3
US-10-076-674-8
; Sequence 8, Application US/10076674
; Publication No. US20030165478A1
; GENERAL INFORMATION:
; APPLICANT: Sokoll, Kenneth K.
; TITLE OF INVENTION: Stabilized Synthetic Immunogen Delivery System
; FILE REFERENCE: Immunogen Delivery System
; CURRENT APPLICATION NUMBER: US/10/076,674
; CURRENT FILING DATE: 2002-04-23
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 8
; LENGTH: 45
; TYPE: PRT
; ORGANISM: Human
US-10-076-674-8

Query Match
Best Local Similarity 64.6%; Score 157; DB 14; Length 45;
Matches 30; Conservative 3; Mismatches 12; Indels 0; Gaps 0;

QY 1 TAAKSKFPSTATYQFGGLSEIKGIVYHRLGVGGHMSYGLRPG 45
DB 1 TAAKSKFPSTATYQFGGLSEIKGIVYHRLGVGGHMSYGLRPG 45

RESULT 4
US-10-355-161A-8
; Sequence 8, Application US/10355161A
; Publication No. US20040009897A1
; GENERAL INFORMATION:
; APPLICANT: Sokoll, Kenneth K.
; TITLE OF INVENTION: Stabilized Synthetic Immunogen Delivery System
; FILE REFERENCE: Immunogen Delivery System
; CURRENT APPLICATION NUMBER: US/10/355,161A
; PRIOR FILING DATE: 2003-01-31
; PRIOR APPLICATION NUMBER: US 10/076674
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 8
; LENGTH: 45
; TYPE: PRT
; ORGANISM: Human
US-10-355-161A-8

Query Match
Best Local Similarity 64.6%; Score 157; DB 15; Length 45;
Matches 30; Conservative 3; Mismatches 12; Indels 0; Gaps 0;

QY 1 TAAKSKFPSTATYQFGGLSEIKGIVYHRLGVGGHMSYGLRPG 45
DB 1 TAAKSKFPSTATYQFGGLSEIKGIVYHRLGVGGHMSYGLRPG 45

DB 1 TAAKSKFPSTATYQFGGLSEIKGIVYHRLGVGGHMSYGLRPG 45

RESULT 5
US-09-848-834A-9
; Sequence 9, Application US/09848834A
; Patent No. US20020076416A1
; GENERAL INFORMATION:
; APPLICANT: Apton Corporation
; TITLE OF INVENTION: Chimeric Peptide Immunogens
; FILE REFERENCE: 1102865-0047
; CURRENT APPLICATION NUMBER: US/09/848,834A
; CURRENT FILING DATE: 2001-05-04
; PRIOR APPLICATION NUMBER: 60/202,328
; PRIOR FILING DATE: 2000-05-05
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 9
; LENGTH: 31
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Chimeric peptide made up of amino acid sequence 288-302 of the Measles virus fusion protein, F linked by a spacer peptide to amino acid sequence 2-10 of the GnRH hormone
; NAME/KEY: MOD RES
; LOCATION: (1)..(11)
; OTHER INFORMATION: Amidated Lysine
; NAME/KEY: PEPTIDE
; LOCATION: (1)..(15)
; OTHER INFORMATION: Peptide corresponds to the amino acid sequences 288-302 of the Measles virus fusion protein, F
; NAME/KEY: PEPTIDE
; LOCATION: (19)..(22)
; OTHER INFORMATION: Spacer peptide
; NAME/KEY: PEPTIDE
; LOCATION: (23)..(31)
; OTHER INFORMATION: Peptide corresponds to amino acid sequences 2-10 of the human GnRH hormone
; NAME/KEY: MOD RES
; LOCATION: (31)..(31)
; OTHER INFORMATION: Amidated glycine or glycylamide
US-09-848-834A-9

Query Match
Best Local Similarity 47.7%; Score 116; DB 9; Length 31;
Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

QY 19 LSEIKGIVYHRLGVGGHMSYGLRPG 44
DB 3 LSEIKGIVYHRLGVGGHMSYGLRPG 30

RESULT 6
US-09-848-834A-17
; Sequence 17, Application US/09848834A
; Patent No. US20020076416A1
; GENERAL INFORMATION:
; APPLICANT: Apton Corporation
; TITLE OF INVENTION: Chimeric Peptide Immunogens
; FILE REFERENCE: 1102865-0047
; CURRENT APPLICATION NUMBER: US/09/848,834A
; CURRENT FILING DATE: 2001-05-04
; PRIOR APPLICATION NUMBER: 60/202,328
; PRIOR FILING DATE: 2000-05-05
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 17
; LENGTH: 47
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Chimeric peptide consisting of amino acid sequence 1-10 of the GnRH hormone
US-09-848-834A-17

Query Match
Best Local Similarity 47.7%; Score 116; DB 9; Length 31;
Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

OTHER INFORMATION: RH hormone linked by a spacer to amino acid sequence 288-302 of
OTHER INFORMATION: the Measles virus protein F linked by a spacer to amino acid seq
OTHER INFORMATION: uence 2-10 of the GnRH hormone
NAME/KEY: MOD_RES
LOCATION: (1)..(1)
OTHER INFORMATION: Pyroglutamic acid or 5-oxoproline
NAME/KEY: MOD_RES
LOCATION: (47)..(47)
OTHER INFORMATION: Amidated-glycine or glycylamide
NAME/KEY: PEPTIDE
LOCATION: (1)..(10)
OTHER INFORMATION: Amino acid sequence 1-10 of the human GnRH hormone
NAME/KEY: PEPTIDE
LOCATION: (11)..(18)
OTHER INFORMATION: Spacer peptide
NAME/KEY: PEPTIDE
LOCATION: (19)..(34)
OTHER INFORMATION: Amino acid sequence 288-302 of the Measles virus fusion protein,
NAME/KEY: PEPTIDE
LOCATION: (35)..(38)
OTHER INFORMATION: Spacer peptide
NAME/KEY: PEPTIDE
LOCATION: (39)..(47)
OTHER INFORMATION: Amino acid sequence 2-10 of the human GnRH hormone
US-09-848-834A-17

Query Match 47.7%; Score 116; DB 9; Length 47;
Best Local Similarity 85.7%; Pred. No. 1.6e-08;
Matches 24; Conservative 0; Mismatches 2; Indels 2; Gaps 1;

QY 19 LSEIKGIVHRLGVGE--HWSYGLRP 44
DB 19 LSEIKGIVHRLGVGEPSLHWSYGLRP 46

RESULT 7
US-09-747-802-72
Sequence 72, Application US/09747802
Publication No. US2003002979A1
GENERAL INFORMATION:
APPLICANT: MANG, CHANG YI
TITLE OF INVENTION: SYNTHETIC PEPTIDE COMPOSITION AS IMMUNOGENS FOR
PREVENTION OF URINARY TRACT INFECTION
FILE REFERENCE: 1151-4165
CURRENT APPLICATION NUMBER: US/09/747,802
CURRENT FILING DATE: 2000-12-22
NUMBER OF SEQ ID NOS: 88
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 72
LENGTH: 16
TYPE: PRT
ORGANISM: Yersinia pseudotuberculosis
US-09-747-802-72

Query Match 34.6%; Score 84; DB 10; Length 16;
Best Local Similarity 100.0%; Pred. No. 0.00013;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPPSYTATYQF 16
DB 1 TAKSKKPPSYTATYQF 16

RESULT 8
US-10-296-734-1467
Sequence 1467, Application US/10296734
Publication No. US20040054137A1
GENERAL INFORMATION:
APPLICANT: Thompson, Scott A
APPLICANT: Ramshaw, Ian A
TITLE OF INVENTION: Synthetic molecules and uses therefor
FILE REFERENCE: Savine
CURRENT APPLICATION NUMBER: US/10/296,734

CURRENT FILING DATE: 2003-08-04
PRIOR APPLICATION NUMBER: AU P07761/00
PRIOR FILING DATE: 2000-05-26
NUMBER OF SEQ ID NOS: 1507
SOFTWARE: PatentIn version 3.2
SEQ ID NO 1467
LENGTH: 16
TYPE: PRT
ORGANISM: Artificial
FEATURE:
OTHER INFORMATION: Invasin immunostimulatory domain
US-10-296-734-1467

Query Match 34.6%; Score 84; DB 15; Length 16;
Best Local Similarity 100.0%; Pred. No. 0.00013;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPPSYTATYQF 16
DB 1 TAKSKKPPSYTATYQF 16

RESULT 9
US-10-789-619-72
Sequence 72, Application US/10789619
Publication No. US20040141993A1
GENERAL INFORMATION:
APPLICANT: MANG, CHANG YI
TITLE OF INVENTION: SYNTHETIC PEPTIDE COMPOSITION AS IMMUNOGENS FOR
PREVENTION OF URINARY TRACT INFECTION
FILE REFERENCE: 1151-4165
CURRENT APPLICATION NUMBER: US/10/789,619
CURRENT FILING DATE: 2004-02-27
NUMBER OF SEQ ID NOS: 88
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 72
LENGTH: 16
TYPE: PRT
ORGANISM: Yersinia pseudotuberculosis
US-10-789-619-72

Query Match 34.6%; Score 84; DB 16; Length 16;
Best Local Similarity 100.0%; Pred. No. 0.00013;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPPSYTATYQF 16
DB 1 TAKSKKPPSYTATYQF 16

RESULT 10
US-10-411-544-32
Sequence 32, Application US/10411544
Publication No. US20030232758A1
GENERAL INFORMATION:
APPLICANT: St. George-Hyslop, Peter
APPLICANT: McLaurin, Joanne
TITLE OF INVENTION: Immunological Methods and Compositions for the Treatment of Alzh
FILE REFERENCE: 1101547
CURRENT APPLICATION NUMBER: US/10/411,544
CURRENT FILING DATE: 2003-04-10
NUMBER OF SEQ ID NOS: 52
SOFTWARE: PatentIn version 3.1
SEQ ID NO 32
LENGTH: 25
TYPE: PRT
ORGANISM: Artificial sequence
FEATURE:
OTHER INFORMATION: chimeric sequence
US-10-411-544-32

Query Match 34.6%; Score 84; DB 15; Length 25;

Mon Feb 14 09:38:20 2005

us-10-076-674a-9.rapb

Best Local Similarity 100.0%; Pred. No. 0.00021;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 19 LSEIKGVYVHRLGVG 35
Db 1 LSEIKGVYVHRLGVG 17

RESULT 11
US-10-351-641-505
Sequence 505, Application US/10351641
GENERAL INFORMATION: US20030186874A1
APPLICANT: Barney, S.
APPLICANT: Merutka, G.
APPLICANT: Anwer, G.
TITLE OF INVENTION: Lamber, M.
TITLE OF INVENTION: HYBRID POLYPEPTIDES WITH ENHANCED PHARMACOKINETIC
FILE REFERENCE: 7872-100
CURRENT FILING DATE: 2003-01-24
PRIOR FILING DATE: 1999-07-09
PRIOR APPLICATION NUMBER: 09/350,641
PRIOR FILING DATE: 1999-07-09
PRIOR APPLICATION NUMBER: 09/315,304
NUMBER OF SEQ ID NOS: 09/082,279
SOFTWARE: FASTSEQ for Windows Version 3.0
SEQ ID NO: 505
LENGTH: 36
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE: OTHER INFORMATION: Core polypeptide
US-10-351-641-505

Query Match
Best Local Similarity 32.5%; Score 79; DB 14; Length 36;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;
Qy 19 LSEIKGVYVHRLGVG---GGEHW 38
Db 13 LSEIKGVYVHRLGVGYNIGSQEW 36

RESULT 12
US-10-267-682-105
Sequence 105, Application US/10267682
GENERAL INFORMATION: US2004003235A1
APPLICANT: Bolognesi, Dani P.
Matthews, Thomas J.
Wild, Carl T.
Barney, Shawn O.
Lambert, Dennis M.
Pettey, Stephen R.
Langlois, Stephen R.
TITLE OF INVENTION: MEMBRANE FUSION-ASSOCIATED EVENTS, INCLUDING HIV
COMPOSITIONS FOR INHIBITION OF
MEMBRANE FUSION-ASSOCIATED EVENTS, INCLUDING HIV
TRANSMISSION
NUMBER OF SEQUENCES: 239
CORRESPONDENCE ADDRESS: 239
ADDRESS: Pennie & Edmonds
CITY: 1155 Avenue of the Americas
STATE: New York
COUNTRY: USA
ZIP: 10036-2711
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
FILING DATE: 08-Oct-2002
CLASSIFICATION: US/10/267,682
PRIOR APPLICATION DATA:
APPLICATION NUMBER: <Unknown>
FILING DATE: 07-JUN-1995
ATTORNEY/AGENT INFORMATION:
NAME: Coruzzi, Laura A.
REGISTRATION NUMBER: 30,742
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 750-9090
TELEFAX: (212) 869-9090
INFORMATION FOR SEQ ID NO: 105:
SEQUENCE CHARACTERISTICS:
LENGTH: 438 amino acids
TYPE: amino acid
STRAND: unknown
TOPOLOGY: unknown
MOLECULE TYPE: protein
SEQUENCE DESCRIPTION: SEQ ID NO: 105:
US-10-267-682-105

Query Match
Best Local Similarity 32.5%; Score 79; DB 15; Length 438;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;
Qy 19 LSEIKGVYVHRLGVG---GGEHW 38
Db 176 LSEIKGVYVHRLGVGYNIGSQEW 199

RESULT 13
US-10-267-748-105
Sequence 105, Application US/10267748
GENERAL INFORMATION: US20040052820A1
APPLICANT: Bolognesi, Dani P.
Matthews, Thomas J.
Wild, Carl T.
Barney, Shawn O.
Lambert, Dennis M.
Pettey, Stephen R.
Langlois, Stephen R.
TITLE OF INVENTION: MEMBRANE FUSION-ASSOCIATED EVENTS, INCLUDING HIV
COMPOSITIONS FOR INHIBITION OF
MEMBRANE FUSION-ASSOCIATED EVENTS, INCLUDING HIV
TRANSMISSION
NUMBER OF SEQUENCES: 239
CORRESPONDENCE ADDRESS: 239
ADDRESS: Pennie & Edmonds
CITY: 1155 Avenue of the Americas
STATE: New York
COUNTRY: USA
ZIP: 10036-2711
COMPUTER READABLE FORM:
MEDIUM TYPE: floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent in Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
FILING DATE: 08-Oct-2002
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/10/267,748
FILING DATE: 07-JUN-1995
ATTORNEY/AGENT INFORMATION

NAME: Coruzzi, Laura A.
 REGISTRATION NUMBER: 30,742
 REFERENCE/DOCKET NUMBER: 7872-029
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (212) 790-9090
 TELEFAX: (212) 869-9741/8864
 TELEX: 66141 PENNIE
 INFORMATION FOR SEQ ID NO: 105:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 438 amino acids
 TYPE: amino acid
 STRANDEDNESS: <Unknown>
 TOPOLOGY: unknown
 MOLECULE TYPE: protein
 SEQUENCE DESCRIPTION: SEQ ID NO: 105:
 US-10-267-748-105

Query Match 32.5%; Score 79; DB 9; Length 438;
 Best Local Similarity 70.8%; Pred. No. 0.029;
 Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGVIYHRLGV---GGEHW 38
 Db 176 LSEIKGVIYHRLGVSYNIGSQEW 199

RESULT 14
 US-09-873-233A-18
 Sequence 18, Application US/09873233A
 Patent No. US20020146434A1
 GENERAL INFORMATION:
 APPLICANT: UEDA, Shigeharu
 APPLICANT: WATANABE, Michiko
 APPLICANT: KAWANISHI, Hitomi
 TITLE OF INVENTION: GENE CODING FOR THE MEASLES VIRUS MUTANT ANTIGEN
 FILE REFERENCE: 0216-0451P
 CURRENT APPLICATION NUMBER: US/09/873,233A
 CURRENT FILING DATE: 2001-06-05
 NUMBER OF SEQ ID NOS: 32
 SOFTWARE: PatentIn Ver. 2.1
 SEQ ID NO 18
 LENGTH: 550
 TYPE: PRT
 ORGANISM: Measles virus
 FEATURE:
 NAME/KEY: UNSURE
 LOCATION: (1)..(550)
 OTHER INFORMATION: any n or Xaa = Unknown
 US-09-873-233A-18

Query Match 32.5%; Score 79; DB 9; Length 550;
 Best Local Similarity 70.8%; Pred. No. 0.038;
 Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGVIYHRLGV---GGEHW 38
 Db 288 LSEIKGVIYHRLGVSYNIGSQEW 311

RESULT 15
 US-09-873-233A-20
 Sequence 20, Application US/09873233A
 Patent No. US20020146434A1
 GENERAL INFORMATION:
 APPLICANT: UEDA, Shigeharu
 APPLICANT: WATANABE, Michiko
 APPLICANT: KAWANISHI, Hitomi
 TITLE OF INVENTION: GENE CODING FOR THE MEASLES VIRUS MUTANT ANTIGEN
 FILE REFERENCE: 0216-0451P
 CURRENT APPLICATION NUMBER: US/09/873,233A
 CURRENT FILING DATE: 2001-06-05
 NUMBER OF SEQ ID NOS: 32
 SOFTWARE: PatentIn Ver. 2.1

SEQ ID NO 20
 LENGTH: 550
 TYPE: PRT
 ORGANISM: Measles virus
 FEATURE:
 NAME/KEY: UNSURE
 LOCATION: (1)..(550)
 OTHER INFORMATION: any n or Xaa = Unknown
 US-09-873-233A-20

Query Match 32.5%; Score 79; DB 9; Length 550;
 Best Local Similarity 70.8%; Pred. No. 0.038;
 Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGVIYHRLGV---GGEHW 38
 Db 288 LSEIKGVIYHRLGVSYNIGSQEW 311

Search completed: February 8, 2005, 17:33:48
 Job time: 128 secs

RESULT 2
Q56889 PRELIMINARY; PRT; 835 AA.
AC O56889
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DE 01-MAR-2004 (TREMBLrel. 26, Last annotation update)
GN Name=InvA;
OS Versinia enterocolitica;
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Bacteriia; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OK NCBI_TaxID=630;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=M1024;
RX MEDLINE=94195100; PubMed=7511772;
RA Pepe J.C., Badger J.L., Miller V.L.;
RT "Growth phase and low pH affect the thermal regulation of the Versinia
RT enterocolitica inv gene.";
RT Mol. Microbiol. 11:123-135(1994).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=M1024;
RX MEDLINE=98046470; PubMed=9387224;
RA Fauconier A., Allacou A., Campos A., Van Elsen A., Cornelle G.R.,
RA Bollen A.;
RT "Flagellar fliA, fliB and fliH genes, organized in an operon, cluster
RT upstream from the inv locus in Versinia enterocolitica.";
RT Microbiology 143:3461-3471(1997).
DR EMBL; Z48169; CA88188.1; -.
DR PIR; S54216; S54216.
DR HSSP; P11922; 1CWV.
DR GO; GO:0007155; P-cell adhesion; IEA.
DR InterPro; IPR003344; Big_1.
DR InterPro; IPR003535; Intimin.
DR Pfam; PF02369; Big_1; 1.
DR PRINTS; PR01369; Intimin.
SQ SEQUENCE 835 AA; 91367 MW; C0176D7766184E3 CRC64;
Query Match 34.6%; Score 84; DB 2; Length 835;
Best Local Similarity 100.0%; Pred. No. 0.029;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 TAKSKKPSYATYQF 16
DB 718 TAKSKKPSYATYQF 733
RESULT 3
VGLF_RINDB STANDARD; PRT; 546 AA.
AC P41360;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DE 05-JUL-2004 (Rel. 44, Last annotation update)
DE Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2;
DE Fusion glycoprotein F1].
GN Name=F.
OS Rinderpest virus (strain RBT1) (RDV).
OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
OC Paramyxoviridae; Paramyxovirinae; Morbillivirinae;
OK NCBI_TaxID=39007;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=M1024;
RX MEDLINE=95088609; PubMed=7996154;
RA Evans S.A., Baron M.D., Chamberlain R.W., Goatsley L., Barrett T.,
RT "Nucleotide sequence comparisons of the fusion protein gene from
RT virulent and attenuated strains of rinderpest virus.";
RT J. Gen. Virol. 75:3611-3617(1994).
CC -1- FUNCTION: This protein directs fusion of viral and cellular

CC membranes.
CC -1- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked.
CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
CC family.
CC
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CC -----
DR EMBL; Z31656; CA83482.1; -.
DR PIR; S47300; S47300.
DR HSSP; P04849; 1SVF.
DR InterPro; IPR000776; Fusion gly.
DR Pfam; PF00523; Fusion gly. F1.
KM Envelope protein; Fusion protein; Glycoprotein; Signal; Transmembrane.
FT SIGNAL 19
FT CHAIN 20 546
FT CHAIN 20 546
FT CHAIN 109 546
FT CHAIN 104 108
FT TRANSSEM 109 133
FT TRANSSEM 484 513
FT DOMAIN 514 517
FT DISULFID 64 191
FT CARBOHYD 25 25
FT CARBOHYD 57 57
FT CARBOHYD 63 63
FT CARBOHYD 518 518
SQ SEQUENCE 546 AA; 58418 MW; 38B539B89344F401 CRC64;
Query Match 32.9%; Score 80; DB 1; Length 546;
Best Local Similarity 63.0%; Pred. No. 0.064;
Matches 17; Conservative 2; Mismatches 4; Indels 4; Gaps 1;
QY 16 FGSLSEIKGIVHRLGVS---GGEHW 38
DB 281 YPSLSEIKGIVHRLGVSYNIGSQEW 307
RESULT 4
Q91HA5 PRELIMINARY; PRT; 546 AA.
ID Q91HA5
AC Q91HA5
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DE 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
DE Fusion protein.
GN Rinderpest virus.
OS Viruses; ssRNA negative-strand viruses; Mononegavirales;
OC Paramyxoviridae; Paramyxovirinae; Morbillivirinae;
OK NCBI_TaxID=11241;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=M1024;
RX MEDLINE=21014265; PubMed=1186456;
RA Ahsan P.K., Sminev A.G., Bezborodova S.V., Starov S.K., Drygin V.V.,
RT "Primary structure of the F-gene from Rinderpest virus strain K.";
RT Mol. Genet. Microbiol. Virol. 4:29-33(2000).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=M1024;
RX MEDLINE=95088609; PubMed=7996154;
RA Evans S.A., Baron M.D., Chamberlain R.W., Goatsley L., Barrett T.,
RT "Nucleotide sequence comparisons of the fusion protein gene from
RT virulent and attenuated strains of rinderpest virus.";
RT J. Gen. Virol. 75:3611-3617(1994).
CC -1- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked (By
CC similarity).
CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
CC family.


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DR EMBL: AY035887; AKK63190.1; -.
DR PIR: P00866; P00866.
DR PIR: P00867; P00867.
DR PIR: P00873; P00873.
DR HSSP: P04849; 1SVF.
DR GO: GO:0019031; C:Viral envelope; IEA.
DR GO: GO:0006948; P:Viral-induced cell-cell fusion; IEA.
DR Pfam: PF00523; Fusion_gly_1.
KM Envelope protein; Fusion protein.
SQ SEQUENCE 546 AA; 58571 MW; 449B2BD7405F0B CRC64;

Query Match 32.5%; Score 79; DB 2; Length 534;
Best Local Similarity 70.8%; Pred. No. 0.064;
Matches 17; Conservative 2; Mismatches 4; Indels 4; Gaps 1;

OY 16 FGLSEIKGVIHRLGCV---GGEHW 38
DB 281 YPSISEIKGVIHRLGCVSYNIGSQEW 307

RESULT 5
OY 076R52 PRELIMINARY; PRT; 531 AA.
AC 076R52;
DT 05-JUL-2004 (TREMBlrel. 27, Created)
DT 05-JUL-2004 (TREMBlrel. 27, Last sequence update)
DE 05-JUL-2004 (TREMBlrel. 27, Last annotation update)
DE Fusion protein.
OS Measles virus.
OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
OC Paramyxoviridae; Paramyxovirinae; Morbillivirinae.
OC NCBI_TaxID=11234;
RN [1]
RP SEQUENCE FROM N.A.
RA Cattaneo R., Billeter M.A.;
RT "Mutated and hypermutated genes of persistent measles viruses which
RT caused lethal human brain diseases.";
RL Virology 0:0-0(0).
CC -1- SUBUNIT: Heterodimer of P1 and P2; disulfide-linked (By
CC similarity).
CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
CC family.
DR EMBL: X16568; CAA34582.1; -.
DR GO: GO:0019031; C:Viral envelope; IEA.
DR GO: GO:0006948; P:Viral-induced cell-cell fusion; IEA.
DR InterPro: IPR000776; Fusion_gly_1.
DR Pfam: PF00523; Fusion_gly_1.
KM Envelope protein; Fusion protein.
SQ SEQUENCE 531 AA; 57568 MW; AF0F45F7AD80DD3 CRC64;

Query Match 32.5%; Score 79; DB 2; Length 531;
Best Local Similarity 70.8%; Pred. No. 0.085;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

OY 19 LSEIKGVIHRLGCV---GGEHW 38
DB 288 LSEIKGVIHRLGCVSYNIGSQEW 311

RESULT 6
VGF_MEASY
ID VGF_MEASY STANDARD; PRT; 534 AA.
AC P26032;
DT 01-MAY-1992 (Rel. 22, Created)
DT 01-MAY-1992 (Rel. 22, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Fusion glycoprotein precursor [Contains: Fusion glycoprotein P2;
DE Fusion glycoprotein P1].
GN Name=;
OS Measles virus (strain Yamagata-1) (Subacute sclerosing panencephalitis
OS virus).
OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
OC Paramyxoviridae; Paramyxovirinae; Morbillivirinae.

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OX NCBI_TaxID=11239;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90385702; PubMed=1698327;
RA Komase K., Haga T., Yoshikawa Y., Sato T.A., Yamaguchi K.;
RT "Molecular analysis of structural protein genes of the Yamagata-1
RT strain of defective subacute sclerosing panencephalitis virus. IV.
RT Nucleotide sequence of the fusion gene.";
RL Virus Genes 4:173-181 (1990).
CC -1- FUNCTION: This protein directs fusion of viral and cellular
CC membranes.
CC -1- SUBUNIT: Heterodimer of P1 and P2; disulfide-linked.
CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
CC family.
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CC or send an email to license@sib-sib.ch).
CC -----
DR EMBL: D10548; BAA01405.1; -.
DR HSSP: P04849; 1SVF.
DR InterPro: IPR000776; Fusion_gly_1.
DR Pfam: PF00523; Fusion_gly_1.
KM Envelope protein; Fusion protein; Glycoprotein; Signal; Transmembrane.
FT SIGNAL 1 23
FT CHAIN 24 534 Fusion glycoprotein P0.
FT CHAIN 24 112 Fusion glycoprotein P2.
FT CHAIN 113 534 Fusion glycoprotein P1.
FT TRANSMEM 113 136 Potential.
FT DOMAIN 137 494 Extracellular (Potential).
FT TRANSMEM 495 515 Potential.
FT DOMAIN 516 534 Cytoplasmic (Potential).
FT DISULFID 68 195 Linkage between P2 and P1 (Potential).
FT CARBOHYD 29 29 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 61 61 N-linked (GlcNAc...) (Potential).
FT CARBOHYD 67 67 N-linked (GlcNAc...) (Potential).
SQ SEQUENCE 534 AA; 57963 MW; F5B21757B643844D CRC64;

Query Match 32.5%; Score 79; DB 1; Length 534;
Best Local Similarity 70.8%; Pred. No. 0.085;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

OY 19 LSEIKGVIHRLGCV---GGEHW 38
DB 288 LSEIKGVIHRLGCVSYNIGSQEW 311

RESULT 7
ID 004243 PRELIMINARY; PRT; 534 AA.
AC 004243;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DT 01-JUN-2003 (TREMBlrel. 24, Last annotation update)
DE Fusion protein.
OS Measles virus.
OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
OC Paramyxoviridae; Paramyxovirinae; Morbillivirinae.
OC NCBI_TaxID=11234;
RN [1]
RP SEQUENCE FROM N.A.
RA Cattaneo R., Billeter M.A.;
RT "Mutated and hypermutated genes of persistent measles viruses which
RT caused lethal human brain diseases.";
RL Virology 0:0-0(0).
CC -1- SUBUNIT: Heterodimer of P1 and P2; disulfide-linked (By
CC similarity).
CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
CC family.

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DR EMBL; X16568; CAA34581.1; -.
DR HSSP; P04849; ISVF.
DR GO; GO:0019031; C:viral envelope; IEA.
DR GO; GO:006948; P:viral-induced cell-cell fusion; IEA.
DR InterPro; IPR00776; Fusion_gly; IEA.
DR pfam; PF00523; Fusion_gly; 1.
DR KX Envelope protein; Fusion protein.
SQ SEQUENCE 534 AA; 57900 MW; 637245E2B5BE044 CRC64;

Query Match      32.5%; Score 79; DB 2; Length 534;
Best Local Similarity 70.8%; Pred.No. 0.085;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY    19 LSEIKGVIVHRLGGV-----GGEHW 38
       |||||
Db     291 LSEIKGVIVHRLGGVSYNIGSQEW 314
```

RESULT 8	Q76RS3	PRELIMINARY;	PRT;	534 AA.
ID	Q76RS3			
AC	Q76RS3;			
DT	05-JUL-2004	(TREMBLrel. 27,		
DT	05-JUL-2004	(TREMBLrel. 27,		
DT	05-JUL-2004	(TREMBLrel. 27,		
DE	Fusion protein.			
OS	Measles virus.			
OC	Viruses; ssRNA negative-strand viruses; Mononegavirales;			
OC	Paramyxoviridae; Paramyxovirinae; Morbillivirinae.			
OX	NCBI_TaxID=11234;			
RN	[1]			
RA	SEQUENCE FROM N.A.			
RA	Cattaneo R., Billerter M.A.;			
RT	"Mutated and hypermutated genes of persistent measles viruses which			
RT	caused lethal human brain diseases.";			
RL	Virology 0:0-0(0).			
CC	-I- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked (By			
CC	similarity).			
CC	-I- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein			
CC	family.			
DR	EMBL; X15567; CAA34575.1; "			
DR	GO; GO:0019031; C:Viral envelope; IEA.			
DR	GO; GO:0006948; P:Viral-induced cell-cell fusion; IEA.			
DR	InterPro; IPR000776; Fusion_gly.			
DR	Pfam; PF00523; Fusion_gly; 1.			
KM	Envelope protein; Fusion_gly; 1.			
SEQUENCE	534 AA; 57944 MW; 10DA7D1A10978B90 CRC64;			

RESULT	9			
ID	004242	PRELIMINARY;	PRT;	537 AA.
AC	004242;			
DT	01-NOV-1996 (TrEMBLrel. 01, Created)			
DT	01-NOV-1996 (TrEMBLrel. 01, Last sequence update)			
DT	01-JUN-2003 (TrEMBLrel. 24, Last annotation update)			
OS	Poison protein.			
DS	Measles virus.			
OC	Viruses; ssRNA negative-strand viruses; Mononegavirales;			
OC	Paramyxoviridae; Paramyxovirine; Morbillivirus.			
OX	NCBI_TaxID=11234;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RA	Cattaneo R., Biller M.A.;			
RT	"Mutated and hypermutated genes of persistent measles viruses which			

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RT caused lethal human brain diseases.",
RL Virology 0:0-0(0) .
CC -1 SUBUNIT: Heterodimer of F1 and F2; disulfide-linked (By
CC similarity) .
CC -1 SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
CC family.
DR EMBL; X1567; CAA34574.1; -.
DR HSSB; P04849; ISVP.
DR GO; GO:0019031; C:viral envelope; IEA.
DR GO; GO:0006948; P:viral-induced cell-cell fusion; IEA.
DR InterPro; IPR000776; Fusion_gly.
DR Pfam; PF00523; Fusion_gly; 1.
KW Envelope protein; Fusion protein.
SQ SEQUENCE 537 AA; 58275 MW; D0A60AC66D979E06 CRC64;

Query Match 32.5%; Score 79; DB 2; Length 537;
Best Local Similarity 70.8%; Pred. NO. 0.086;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGIVVHRLEGV---GGEHW 38
|||||
DQ 291 LSEIKGIVVHRLEGVSYNIGQEW 314
|||||

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RESULT 10
ID O9PX4 PRELIMINARY; PRT; 545 AA.
AC O9PX4;
DT 01-MAY-2000 (TrEMBLrel. 13, Created)
DT 01-MAY-2000 (TrEMBLrel. 13, last sequence update)
DE 05-JUL-2004 (TrEMBLrel. 27, last annotation update)
DE Fusion protein.
ME Measles virus.
OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.
NCBI_TaxID=11234;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=OSA-3;
RX MEDLINE=22075939; PubMed=12076936; DOI=10.1016/S0168-1702(02)00042-4;
RA Ning X., Ayata M., Kimura M., Komase K., Furukawa K., Seto T., Ito N.
RA Shingai N., Matsumura I., Yamano T., Ogura H.;
RT "Alterations and diversity in the cytoplasmic tail of the fusion
RT protein of subacute sclerosing panencephalitis virus strains isolated
RT in Osaka, Japan.";
RL Virus Res. 86:123-131(2002).
CC -1- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked (By
CC similarity).
CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
CC family.
EMBL: AF179440; AAF02705.1; -.
EMBL: AF179439; AAF02704.1; -.
DR HSBP; P04849; ISVF.
DR GO; GO:0019031; C:viral envelope; IEA.
DR GO; GO:0006948; P:viral-induced cell-cell fusion; IEA.
DR InterPro; IPR000776; Fusion_gly.
DR Pfam; PF00523; Fusion_gly; I.
KW Envelope protein; Fusion protein.
SQ SEQUENCE 545 AA; 58907 MW; 0234C28AB193E77D CRC64;

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Query Match      32.5%   Score 79; DB 2; Length 545;
Best Local Similarity 70.8%; Pred. No. 0.087;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

Oy      19 LSEIKGVIVHRLEGV---GGEHW 38
          ||||| | | | | | | | | | : |
Db      288 LSEIKGVIVHRLEGVSYNIGSQEW 311

RESULT 11
VGLF_RINDR
AC P41356; STANDARD; PRT; 546 AA.
```

DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-FEB-1995 (Rel. 31, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2;
 DE Fusion glycoprotein F1].
 GN Name=F;
 OS Rinderpest virus (strain RBOK) (RDV).
 OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbilliviruses.
 NCBI_TaxID=36409;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=95088609; PubMed=7996154;
 RA Evans S.A., Baron M.D., Chamberlain R.W., Goateley L., Barrett T.;
 RT "Nucleotide sequence comparisons of the fusion protein gene from
 RT virulent and attenuated strains of rinderpest virus";
 RL J. Gen. Virol. 75:3611-3617(1994).
 CC -1- FUNCTION: This protein directs fusion of viral and cellular
 CC membranes.
 CC -1- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked.
 CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
 CC family.
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 CC -----
 DR EMBL: Z30700; CAA83186.1; -;
 DR EMBL: Z30697; CAA83181.1; -;
 DR PIR: S47305; S47305.
 DR HSSP: P04849; 1SVF.
 DR InterPro: IPR000776; Fusion gly.
 DR Pfam: PF00523; Fusion gly. 1.
 KM Envelope protein; Fusion protein; Glycoprotein; Signal; Transmembrane.
 FT SIGNAL 1
 FT CHAIN 19
 FT CHAIN 20 546 Fusion glycoprotein F0.
 FT CHAIN 20 108 Fusion glycoprotein F2.
 FT CHAIN 109 546 Fusion glycoprotein F1.
 FT DOMAIN 104 108 Arg/Lys-rich (basic).
 FT TRANSMEM 109 133 Potential.
 FT TRANSMEM 484 513 Potential.
 FT DOMAIN 514 517 Arg/Lys-rich (basic).
 FT DISULFID 64 191 Linkage between F2 and F1 (Potential).
 FT CARBOHYD 25 25 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 57 57 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 63 63 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 518 518 N-linked (GlcNAc...) (Potential).
 SQ SEQUENCE 546 AA; 58705 MW; ED3DF8AFDEBCB95 CRC64;
 Query Match 32.5%; Score 79; DB 1; Length 546;
 Best Local Similarity 59.3%; Pred. No. 0.087; Mismatches 4; Indels 4; Gaps 1;
 Matches 16; Conservative 3; Mismatches 4; Indels 4; Gaps 1;
 QY 16 FGGLSEIKGYIVHRLGCV---GGEHW 38
 DB 281 YPSLSEIKGYIVHRLGCVSYNIGSQEW 307
 RESULT 12
 VGLF MEASA STANDARD; PRT; 550 AA.
 AC P35973;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2;
 DE Fusion glycoprotein F1].
 GN Name=F;
 OS Measles virus (strain AIR-C) (Subacute sclerosing panencephalitis

OS virus).
 OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbilliviruses.
 NCBI_TaxID=36408;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93227570; PubMed=8470368;
 RA Mori T., Saeki K., Hashimoto H., Makino S.;
 RT "Molecular cloning and complete nucleotide sequence of genomic RNA of
 RT the AIR-C strain of attenuated measles virus";
 RL Virus Genes 7:67-81(1993).
 CC -1- FUNCTION: This protein directs fusion of viral and cellular
 CC membranes.
 CC -1- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked.
 CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
 CC family.
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 CC -----
 DR EMBL: S58435; AAB26145.1; -;
 DR PIR: E48556; E48556.
 DR HSSP: P04849; 1SVF.
 DR InterPro: IPR000776; Fusion gly.
 DR Pfam: PF00523; Fusion gly. 1.
 KM Envelope protein; Fusion protein; Glycoprotein; Signal; Transmembrane.
 FT SIGNAL 23
 FT CHAIN 24 550 Fusion glycoprotein F0.
 FT CHAIN 24 112 Fusion glycoprotein F2.
 FT CHAIN 113 550 Fusion glycoprotein F1.
 FT TRANSMEM 113 136 Potential.
 FT DOMAIN 137 494 Extracellular (Potential).
 FT TRANSMEM 495 515 Potential.
 FT DOMAIN 516 550 Cytoplasmic (Potential).
 FT DISULFID 68 190 Linkage between F2 and F1 (Potential).
 FT CARBOHYD 29 29 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 61 61 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 67 67 N-linked (GlcNAc...) (Potential).
 SQ SEQUENCE 550 AA; 59540 MW; AAC4DAB92DEBD938 CRC64;
 Query Match 32.5%; Score 79; DB 1; Length 550;
 Best Local Similarity 70.8%; Pred. No. 0.088; Mismatches 2; Indels 4; Gaps 1;
 Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;
 QY 19 LSEIKGYIVHRLGCV---GGEHW 38
 DB 288 LSEIKGYIVHRLGCVSYNIGSQEW 311
 RESULT 13
 VGLF MEASE STANDARD; PRT; 550 AA.
 AC P08300;
 DT 01-AUG-1988 (Rel. 08, Created)
 DT 01-AUG-1988 (Rel. 08, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Fusion glycoprotein precursor [Contains: Fusion glycoprotein F2;
 DE Fusion glycoprotein F1].
 GN Name=F;
 OS Measles virus (strain Edmonston) (Subacute sclerosing panencephalitis
 OS virus).
 OS Measles virus (strain Halle) (Subacute sclerosing panencephalitis
 OS virus).
 OS Measles virus (strain Leningrad-16) (Subacute sclerosing panencephalitis
 OS virus).
 OS Measles virus (strain Edmonston-Zagreb) (Subacute sclerosing
 OS panencephalitis virus).
 OS Measles virus (strain Philadelphia-26) (Subacute sclerosing

OS panencephalitis virus), and
 OS Measles virus (strain Edmonston B) (Subacute sclerose panencephalitis
 OS virus).
 OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.
 OX NCBI_TaxID=11235, 11236, 70147, 70149, 70148, 70146;
 [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Edmonston;
 RX MEDLINE=87071668; PubMed=3788062;
 RA Richardson C.D., Hull D., Greer P., Haesel K., Berkovich A.,
 RA Englund G., Bellini W.J., Rima B., Lazzarini R.A.;
 RT "The nucleotide sequence of the mRNA encoding the fusion protein of
 RT measles virus (Edmonston strain): a comparison of fusion proteins from
 RT several different paramyxoviruses.";
 RL Virology 155:508-523 (1986).
 RN [2]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Hallie;
 RX MEDLINE=87224816; PubMed=3585281;
 RA Buckland R., Gerald C., Barker R., Wild T.F.;
 RT "Fusion glycoprotein of measles virus: nucleotide sequence of the gene
 RT and comparison with other paramyxoviruses.";
 RL J. Gen. Virol. 68:1695-1703 (1987).
 RN [3]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Edmonston;
 RX MEDLINE=90085790; PubMed=2596022;
 RA Cattaneo R., Schmid A., Spielhofer P., Kaelin K., Bacsko K.,
 RA Meulen V., Pardowitz J., Flanagan S., Rima B.K., Udem S.A.;
 RT "Mutated and hypermutated genes of persistent measles viruses which
 RT caused lethal human brain diseases.";
 RL Virology 173:415-425 (1989).
 RN [4]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Edmonston;
 RX MEDLINE=92263801; PubMed=1585658;
 RA Schmid A., Spielhofer P., Cattaneo R., Bacsko K., Ter Meulen V.,
 RA Biller M.A.;
 RT "Subacute sclerosing panencephalitis is typically characterized by
 RT alterations in the fusion protein cytoplasmic domain of the persisting
 RT measles virus.";
 RL Virology 188:910-915 (1992).
 RN [5]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Edmonston, Edmonston-Zagreb, and Leningrad-16;
 RX MEDLINE=94249283; PubMed=8191786; DOI=10.1016/0168-1702(94)90025-6;
 RA Rota J.S., Wang Z.D., Rota P.A., Bellini W.J.;
 RT "Comparison of sequences of the H, F, and N coding genes of measles
 RT virus vaccine strains ";
 RL Virus Res. 31:317-330 (1994).
 RN [6]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Philadelphia-26;
 RX MEDLINE=94303181; PubMed=8030232;
 RA Hummel K.B., Vanchiere J.A., Bellini W.J.;
 RT "Restriction of fusion protein mRNA as a mechanism of measles virus
 RT persistence.";
 RL Virology 202:665-672 (1994).
 RN [7]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Edmonston B;
 RA Biller M.A.;
 RL Submitted (OCT-1995) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: This protein directs fusion of viral and cellular
 CC membranes.
 CC -1- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked.
 CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
 CC family.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
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 CC -----
 CC EMBL; M14915; AAA46423.1; -;
 DR EMBL; X05597; CAA29090.1; ALT_INIT.
 DR EMBL; K01711; AAA75498.1; ALT_INIT.
 DR EMBL; K01711; AAA75499.1; -;
 DR EMBL; U03657; AAA56647.1; ALT_INIT.
 DR EMBL; U03659; AAA56649.1; ALT_INIT.
 DR EMBL; U03670; AAA56660.1; ALT_INIT.
 DR EMBL; U08416; AAA50550.1; ALT_INIT.
 DR EMBL; Z66517; CAA91367.1; ALT_INIT.
 DR EMBL; Z66517; CAA91368.1; -;
 DR HSPSP; P04849; ISVP.
 DR InterPro; IPR000776; Fusion_gly.
 DR Pfam; PF00523; Fusion_gly_1.
 KW Envelope protein; Fusion protein; Glycoprotein; Signal; Transmembrane.
 FT SIGNAL 1 23
 FT CHAIN 24 550 Fusion glycoprotein F0.
 FT CHAIN 24 112 Fusion glycoprotein F2.
 FT CHAIN 113 550 Fusion glycoprotein F1.
 FT TRANSMEM 113 136 Potential.
 FT DOMAIN 137 494 Extracellular (Potential).
 FT TRANSMEM 495 515 Potential.
 FT DOMAIN 516 550 Cytoplasmic (Potential).
 FT DISULFID 68 195 Linkage between F2 and F1 (Potential).
 FT CARBOHYD 29 61 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 61 61 N-linked (GlcNAc...) (Potential).
 FT CARBOHYD 67 67 N-linked (GlcNAc...) (Potential).
 SQ SEQUENCE 550 AA; 59532 MW; 7AAAF1CA82169093 CRC64;
 Query Match 32.5%; Score 79; DB 1; Length 550;
 Best Local Similarity 70.8%; Pred. No. 0.088;
 Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;
 Oy 19 LSEIKGVYHRLBEGV---GGEHW 38
 Db 288 LSEIKGVYHRLBEGVSYNIGSQEW 311
 RESULT 14
 P90330 PRELIMINARY; PRT; 550 AA.
 AC P90330
 DT 01-MAY-1997 (TREMBLrel. 03, Created)
 DT 01-MAY-1997 (TREMBLrel. 03, Last sequence update)
 DT 01-OCT-2003 (TREMBLrel. 25, Last annotation update)
 DE Fusion protein.
 OS Measles virus.
 OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbillivirus.
 OX NCBI_TaxID=11234;
 RN [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Nagahata;
 RA Sheng J., Watanabe M., Ueda S.;
 RL Submitted (AUG-1995) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RN SEQUENCE FROM N.A.
 RC STRAIN=Nagahata;
 RA Sheng J., Watanabe M., Ueda S.;
 RL Submitted (AUG-1995) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked (By
 CC similarity).
 CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
 CC family.
 CC EMBL; D63924; BAA09951.1; -;
 DR PIR; P00376; P00376.
 DR HSPSP; P04849; ISVP.
 DR GO; GO:0019031; C:Viral envelope; IEA.
 DR GO; GO:0006948; P:Viral-induced cell-cell fusion; IEA.

DR InterPro: IPR000776; Fusion gly.
 DR Pfam: PF00523; Fusion gly; 1
 KM Envelope protein; Fusion protein.
 SQ SEQUENCE 550 AA; 59589 MW; 73E7BD457ABA39B7 CRC64;

Query Match 32.5%; Score 79; DB 2; Length 550;
 Best Local Similarity 70.8%; Pred. No. 0.088;
 Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGVIHRLGV---GGEHM 38
 DB 288 LSEIKGVIHRLGVSYNIGSQEW 311

RESULT 15

ID P90331 PRELIMINARY; PRT; 550 AA.
 AC P90331;
 DT 01-MAY-1997 (T-EMBLrel. 03, Created)
 DT 01-MAY-1997 (T-EMBLrel. 03, Last sequence update)
 DT 05-JUL-2004 (T-EMBLrel. 27, Last annotation update)
 DE Fusion protein.
 OS Measles virus.
 OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
 OC Paramyxoviridae; Paramyxovirinae; Morbillivirinae;
 OC NCBI_TaxID=11234;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Nagahata;
 RA Sheng J., Watanabe M., Ueda S.;
 RL Submitted (Aug-1995) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Nagahata;
 RA Sheng J., Nakanishi M., Watanabe M., Ueda S.;
 RL Submitted (Aug-1995) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Nagahata;
 MEDLINE=22072939; PubMed=12076836; DOI=10.1016/S0168-1702(02)00042-4;
 RA Ning X., Ayata M., Kimura M., Komase K., Furukawa K., Seto T., Ito N.,
 RA Shingai M., Matsunaga I., Yamano T., Ogura H.;
 RT "Alterations and diversity in the cytoplasmic tail of the fusion
 protein of subacute sclerosing panencephalitis virus strains isolated
 in Osaka, Japan.";
 RT Virus Res. 86:123-131(2002).
 CC -1- SUBUNIT: Heterodimer of F1 and F2; disulfide-linked (By
 similarity).
 CC -1- SIMILARITY: Belongs to the paramyxoviruses fusion glycoprotein
 family.
 CC EMBL: D63926; BAA09958.1; -.
 DR EMBL: AF179431; AAF02696.1; -.
 DR PIR: P00376; P00376.
 DR HSSP: P04849; 1SVF.
 DR GO: GO:0019031; C:Viral envelope; IEA.
 DR GO: GO:0006948; P:Viral-induced cell fusion; IEA.
 DR InterPro: IPR000776; Fusion gly.
 DR Pfam: PF00523; Fusion gly; 1.
 KM Envelope protein; Fusion protein.
 SQ SEQUENCE 550 AA; 59530 MW; 97C991C7E2169839 CRC64;

Query Match 32.5%; Score 79; DB 2; Length 550;
 Best Local Similarity 70.8%; Pred. No. 0.088;
 Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGVIHRLGV---GGEHM 38
 DB 288 LSEIKGVIHRLGVSYNIGSQEW 311

Search completed: February 8, 2005, 17:30:50
 Job time : 174 secs

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OM protein - protein search, using sw model

Run on: February 8, 2005, 17:19:57 ; Search time 39 Seconds

(without alignments)
111.019 Million cell updates/sec

Title: US-10-076-674A-9

Perfect score: 243

Sequence: 1 TAKSKKFPSTATYQFGSL.....IVHRLGVGGEHWSYGLRFG 45

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PIR 79: *
1: p1r1: *
2: p1r2: *
3: p1r3: *
4: p1r4: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	84	34.6	835	1 S54216	invasin - Yersinia
2	84	34.6	835	2 S11442	invasin - Yersinia
3	80	32.9	546	2 S47300	gene F protein - r
4	79	32.5	282	2 P00376	cell fusion glycop
5	79	32.5	282	2 P00388	cell fusion glycop
6	79	32.5	534	1 J00274	cell fusion glycop
7	79	32.5	546	1 VGNZKX	cell fusion glycop
8	79	32.5	546	1 VGNZKX	cell fusion glycop
9	79	32.5	550	1 S47305	gene F protein - r
10	79	32.5	553	1 VGNZMV	cell fusion glycop
11	74	30.5	90	1 RHMSG	gonadolibetin prec
12	74	30.5	92	1 RHMSG	gonadolibetin prec
13	74	30.5	546	1 VGNZRL	gonadolibetin prec
14	72	29.6	542	2 J02223	cell fusion glycop
15	72	29.6	552	2 S47034	cell fusion glycop
16	72	29.6	631	1 A48346	cell fusion glycop
17	72	29.6	631	1 VGNZPD	cell fusion glycop
18	72	29.6	662	1 VGNZCD	cell fusion glycop
19	72	29.6	662	2 S21382	cell fusion glycop
20	71	29.2	92	1 RHMSG	gonadolibetin prec
21	70	28.8	67	2 I78541	gonadolibetin prec
22	70	28.8	986	2 A29645	invasin - Yersinia
23	67	27.6	546	2 S53386	cell fusion glycop
24	64.5	26.5	265	2 T05668	cell fusion glycop
25	64	26.3	636	2 S47299	pollen allergen ho
26	63.5	26.1	89	2 S15123	gonadolibetin prec
27	61	25.1	900	2 JH0157	cellulase (EC 3.2.
28	60	24.7	10	1 RHMSG	gonadolibetin - p1
29	60	24.7	10	1 RHMSG	gonadolibetin - p1

30	60	24.7	1085	2 T03531	cellulase (EC 3.2.
31	57	23.3	584	2 J01229	transcription init
32	57	23.3	675	2 J39065	gonadolibetin i -
33	56	23.0	10	1 RHAQ1	gonadolibetin i -
34	56	23.0	92	2 I50644	50S ribosomal prot
35	56	23.0	147	2 A84546	ribosomal protein
36	56	23.0	162	2 T49957	hypothetical prote
37	56	23.0	187	2 T47342	hypothetical prote
38	56	23.0	473	2 T38350	threonine-tRNA lig
39	56	23.0	551	2 B64728	yabn protein - Eac
40	56	23.0	552	2 B90638	probable transport
41	56	23.0	552	2 B85489	probable acyl-CoA
42	56	23.0	601	2 D83583	probable glucose-6
43	55.5	22.8	489	2 B69664	hypothetical prote
44	54.5	22.4	263	2 T47536	nitrate reductase
45	54.5	22.4	710	2 B69665	

ALIGNMENTS

RESULT 1

S54216
invasin - Yersinia enterocolitica (strain W1024)

C:Species: Yersinia enterocolitica

A:Variety: strain W1024

C>Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 09-Jul-2004

C/Accession: S54216

R:Fauconier, A.; Allaoui, A.; Van Elsen, A.; Cornelle, G.; Bollen, A.

submitted to the EMBL Data Library, February 1995

A:Description: Clustering of flagellar genes around invA, the Yersinia enterocolitica inv

A:Reference number: S54213

A:Accession: S54216

A:Molecule type: DNA

A:Residues: 1-835 <FAU>

A:Cross-references: UNIPROT:Q56889; EMBL:Z48169; NID:9793891; PIDN:CAA88188.1; PID:G79385

A:Experimental source: strain W1024; serotype O:9

C:Genetics:

A:Gene: invA

C:Superfamily: Invasin

Query Match 34.6%; Score 84; DB 1; Length 835;
Best Local Similarity 100.0%; Pred. No. 0.0097;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKFPSTATYQF 16
|||||
Db 718 TAKSKKFPSTATYQF 733

RESULT 2

S11442

invasin - Yersinia enterocolitica

C:Species: Yersinia enterocolitica

C>Date: 21-Nov-1993 #sequence_revision 10-Nov-1995 #text_change 09-Jul-2004

R:Young, V.B.; Miller, V.L.; Falkow, S.; Schoolnik, G.K.

Mol. Microbiol. 4, 1119-1128, 1990

A:Title: Sequence, localization and function of the invasin protein of Yersinia enterocoli

A:Reference number: S11442; NID:91041720; PMID:2233250

A:Accession: S11442

A:Molecule type: DNA

A:Residues: 1-835 <YOU>

A:Cross-references: UNIPROT:P19196; EMBL:X53368; NID:948573; PIDN:CAA37448.1; PID:G48574

A:Experimental source: strain 8081c

C:Genetics:

A:Gene: invA

C:Superfamily: Invasin

Query Match 34.6%; Score 84; DB 2; Length 835;
Best Local Similarity 100.0%; Pred. No. 0.0097;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TAKSKKPSYATAYOR 16
|||||
Db 718 TAKSKKPSYATAYOR 733

RESULT 3

gene F protein - rinderpest virus
C/Species: rinderpest virus
C/Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 09-Jul-2004
C/Accession: S47300, PQ0865
R/Schulz, S.A.; Baron, M.D.; Chamberlain, R.W.; Goatley, L.; Barrett, T.
submitted to the EMBL Data Library, March 1994
A/Description: The complete nucleotide sequence of the fusion protein gene of the vacci
A/Reference number: S47299
A/Accession: S47300
A/Molecule type: DNA
A/Residues: 1-546 <EVA>
A/Cross-references: UNIPROT:P41360; EMBL:Z31656; NID:G535406; PIDD:CA83482.1; PID:G5354
R/Chamberlain, R.W.; Wamwayi, H.M.; Hockley, E.; Shaila, M.S.; Goatley, L.; Knowles, N.J
J. Gen. Virol. 74, 2775-2780, 1993
A/Title: Evidence for different lineages of rinderpest virus reflecting their geographic
A/Reference number: PQ0865; MUID:94103786; PMID:8277286
A/Accession: PQ0865
A/Molecule type: mRNA
A/Residues: 86-191 <CHA>
C/Genetics:
A/Gene: F
C/Superfamily: paramfluenza virus cell fusion protein
C/Keywords: glycoprotein; membrane fusion; transmembrane protein

Query Match 32.5%; Score 79; DB 2; Length 282;
Best Local Similarity 70.8%; Pred. No. 0.013;
Matches 17; Conservative 2; Mismatches 4; Indels 4; Gaps 1;

QY 16 FGLSEIKGYIVRLRGV----GGEHW 38
:|||||
Db 281 YPSELSEIKGYIVRLRGVSYNIGSQEW 307

RESULT 4

cell fusion glycoprotein - measles virus (strain TT) (fragment)
C/Species: measles virus
C/Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 09-Jul-2004
C/Accession: PQ0376
R/Schulz, T.F.; Hoed, J.G.; Whitby, D.; Tizard, E.J.; Dillon, M.J.; Weiss, R.A.
J. Gen. Virol. 73, 1581-1586, 1992
A/Title: A measles virus isolate from a child with Kawasaki disease: sequence comparison
A/Reference number: PQ0374; MUID:92300360; PMID:1607874
A/Accession: PQ0376
A/Molecule type: genomic RNA
A/Residues: 1-282 <SCH>
A/Cross-references: UNIPROT:Q83529; UNIPROT:Q93055; UNIPROT:Q91C36; UNIPROT:P88973; UNIP
ROT:P90330; UNIPROT:Q9QEW7; UNIPROT:Q9WAK4; UNIPROT:Q83525; UNIPROT:Q8318; UNIPROT:Q89
3521; UNIPROT:Q83530; UNIPROT:Q91248; UNIPROT:Q91Q22; UNIPROT:Q9QEW8; UNIPROT:Q04244
C/Genetics:
A/Gene: F
C/Superfamily: paramfluenza virus cell fusion protein
C/Keywords: glycoprotein; membrane fusion

Query Match 32.5%; Score 79; DB 2; Length 282;
Best Local Similarity 70.8%; Pred. No. 0.013;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGYIVRLRGV----GGEHW 38
|||||
Db 20 LSEIKGYIVRLRGVSYNIGSQEW 43

RESULT 5

PQ0388
cell fusion glycoprotein - measles virus (strain Schwarz vaccine) (fragment)

C/Species: measles virus
C/Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 09-Jul-2004
C/Accession: PQ0388
R/Schulz, T.F.; Hoed, J.G.; Whitby, D.; Tizard, E.J.; Dillon, M.J.; Weiss, R.A.
J. Gen. Virol. 73, 1581-1586, 1992
A/Title: A measles virus isolate from a child with Kawasaki disease: sequence compariso
A/Reference number: PQ0374; MUID:92300360; PMID:1607874
A/Accession: PQ0388
A/Molecule type: genomic RNA
A/Residues: 1-282 <SCH>
A/Cross-references: UNIPROT:Q83525; UNIPROT:Q83530
C/Genetics:
A/Gene: F
C/Superfamily: paramfluenza virus cell fusion protein
C/Keywords: glycoprotein; membrane fusion

Query Match 32.5%; Score 79; DB 2; Length 282;
Best Local Similarity 70.8%; Pred. No. 0.013;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGYIVRLRGV----GGEHW 38
|||||
Db 20 LSEIKGYIVRLRGVSYNIGSQEW 43

RESULT 6

cell fusion glycoprotein precursor - subacute sclerosing panencephalitis virus (strain Y
JUN274
N/Contains: fusion glycoprotein F1; fusion glycoprotein F2
C/Species: subacute sclerosing panencephalitis virus, SSPEV
C/Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 16-Jun-2000
C/Accession: JUN274
R/Komase, K.; Haga, T.; Yoshikawa, Y.; Sato, T.A.; Yamanochi, K.
Virus Genes 4, 173-181, 1990
A/Title: Molecular analysis of structural protein genes of the Yamagata-1 strain of defec
A/Reference number: JUN274; MUID:90385702; PMID:1698327
A/Accession: JUN274
A/Molecule type: mRNA
A/Residues: 1-534 <KOM>
A/Cross-references: EMBL:D10548; NID:G222256; PIDD:BA01405.1; PID:G222257
A/Note: the authors translated the codon GTA for residue 459 as Gly and GCG for residue
C/Genetics:
A/Gene: F
C/Superfamily: paramfluenza virus cell fusion protein
C/Keywords: glycoprotein; membrane fusion; transmembrane protein
F1-22/Domain: signal sequence #status predicted <SIG>
F123-107/Product: cell fusion glycoprotein F2 #status predicted <F2>
F1208-534/Product: cell fusion glycoprotein F1 #status predicted <F1>
F1498-514/Domain: transmembrane #status predicted <TM>
F16,29,61,67/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 32.5%; Score 79; DB 1; Length 534;
Best Local Similarity 70.8%; Pred. No. 0.026;
Matches 17; Conservative 1; Mismatches 2; Indels 4; Gaps 1;

QY 19 LSEIKGYIVRLRGV----GGEHW 38
|||||
Db 288 LSEIKGYIVRLRGVSYNIGSQEW 311

RESULT 7

cell fusion glycoprotein precursor - rinderpest virus (strain Kabete O)
N/Contains: fusion glycoprotein F1; fusion glycoprotein F2
C/Species: rinderpest virus
C/Date: 31-Dec-1989 #sequence_revision 31-Dec-1989 #text_change 09-Jul-2004
C/Accession: A31051
R/Hsu, D.; Yamataka, M.; Miller, J.; Dale, B.; Grubman, M.; Ylma, T.
Virology 166, 149-153, 1988
A/Title: Cloning of the fusion gene of rinderpest virus: comparative sequence analysis w
A/Reference number: A31051; MUID:88322864; PMID:3413983
A/Accession: A31051
A/Molecule type: genomic RNA

R;Maason, A.J.; Hayflick, J.S.; Zoeller, R.T.; Young III, W.S.; Phillips, H.S.; Nikolic, Science 234, 1366-1371, 1986
A;Title: A deletion truncating the gonadotropin-releasing hormone gene is responsible for
A;Reference number: A47578; MUID:87069928; PMID:3024317
A;Accession: A47578
A;Molecule type: DNA
A;Residues: 1-90 <MAS>
A;Cross-references: UNIPROT:P13562; EMBL:M4872; NID:G193576; PID:AAA37717.1; PID:G3871
C;Genetics:
A;Introns: 45/3; 77/3
C;Function:
A;Description: gonadoliberin stimulates pituitary secretion of luteotropin and follitropin
A;Note: gonadoliberin-associated protein may have prolactin release inhibiting activity
C;Keywords: amidated carboxyl end; hormone; hypothalamus; pyroglyutamic acid
F;1-23/Domain: signal sequence #status predicted <SIG>
F;22-31/Product: gonadoliberin #status predicted <GLB>
F;35-90/Product: gonadoliberin-associated protein #status predicted <GAP>
F;32/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status predicted
F;31/Modified site: amidated carboxyl end (Gly) (amide in mature form from following gly

Query Match 30.5%; Score 74; DB 1; Length 90;
Best Local Similarity 75.0%; Pred. No. 0.016;
Matches 12; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 30 LEQVGEHMSYGLRPG 45
DB 16 LEQSSQHMSYGLRPG 31
||| :|||||
| :|

RESULT 12
RHRNG
N;Alternate names: gonadoliberin-associated protein (GAP); gonadotropin releasing hormone
N;Contains: gonadoliberin; prolactin release-inhibiting factor
C;Species: Rattus norvegicus (Norway rat)
C;Date: 31-Mar-1988 #sequence_revision 31-Mar-1988 #text_change 09-Jul-2004
C;Accession: A40147; B26173; A48410
R;Bond, C.T.; Hayflick, J.S.; Seeburg, P.H.; Adelman, J.P.
Mol. Endocrinol. 3, 1257-1262, 1989
A;Title: The rat gonadotropin-releasing hormone: SH locus: structure and hypothalamic ex
A;Reference number: A40147; MUID:8938461; PMID:2476669
A;Accession: A40147
A;Molecule type: DNA
A;Residues: 1-92 <DNA>
A;Cross-references: UNIPROT:P07490; GB:M31670; NID:G204447; PID:AAA41264.1; PID:G204448
R;Adelman, J.P.; Maason, A.J.; Hayflick, J.S.; Seeburg, P.H.
Proc. Natl. Acad. Sci. U.S.A. 83, 179-183, 1986
A;Title: Isolation of the gene and hypothalamic cDNA for the common precursor of gonadot
A;Reference number: A94090; MUID:86094338; PMID:2867548
A;Accession: B26173
A;Molecule type: mRNA
A;Residues: 1-92 <ADEN>
A;Cross-references: GB:M12579; NID:G204445; PID:AAA41263.1; PID:G204446
R;Maier, C.C.; Marchetti, B.; LeBoeuf, R.D.; Blalock, J.E.
Cell. Mol. Neurobiol. 12, 447-454, 1992
A;Title: Thyrocytes express a mRNA that is identical to hypothalamic luteinizing hormone
A;Reference number: A48410; MUID:93105480; PMID:1468115
A;Accession: A48410
A;Status: preliminary
A;Molecule type: mRNA
A;Residues: 1-92 <MAI>
A;Cross-references: GB:S50870; NID:G262059; PID:AAAB2572.1; PID:G262060
A;Experimental source: thymus
A;Note: sequence extracted from NCBI backbone (NCBIN:121082, NCIP:121083)
C;Genetics:
A;Introns: 47/3; 79/3
C;Function:
A;Description: stimulates pituitary secretion of luteotropin and follitropin
A;Note: gonadoliberin-associated protein may have prolactin release inhibiting activity
C;Keywords: amidated carboxyl end; hormone; hypothalamus; placenta; pyroglyutamic acid; r
F;1-23/Domain: signal sequence #status predicted <SIG>

F;24-92/Product: progadoliberin #status predicted <FCN>
F;24-33/Product: gonadoliberin #status predicted <GHN>
F;37-92/Product: prolactin release-inhibiting factor #status predicted <PIF>
F;24/Modified site: pyroglutamate carboxylic acid (Gln) (in mature form) #status predicted
F;33/Modified site: amidated carboxyl end (Gly) (amide in mature form from following gly

Query Match 30.5%; Score 74; DB 1; Length 92;
Best Local Similarity 75.0%; Pred. No. 0.016;
Matches 12; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 30 LEQVGEHMSYGLRPG 45
DB 18 LEQSSQHMSYGLRPG 33
||| :|||||
| :|

RESULT 13
VGNZRL
cell fusion glycoprotein precursor - rinderpest virus (strain L)
N;Contains: fusion glycoprotein F1; fusion glycoprotein F2
C;Species: rinderpest virus
C;Date: 30-Sep-1989 #sequence_revision 30-Sep-1989 #text_change 09-Jul-2004
C;Accession: A28921
R;Tanikawa, K.; Yoshikawa, Y.; Yamanouchi, K.
Virology 164, 523-530, 1988
A;Title: Fusion glycoprotein (F) of rinderpest virus: entire nucleotide sequence of the
A;Reference number: A28921; MUID:88219541; PMID:3285575
A;Accession: A28921
A;Molecule type: mRNA
A;Residues: 1-546 <TSU>
A;Cross-references: UNIPROT:P10864; GB:M20870; NID:G333898; PID:AAA47399.1; PID:G333899
C;Genetics:
A;Gene: F
C;Superfamily: parainfluenza virus cell fusion protein
C;Keywords: glycoprotein; membrane fusion; transmembrane protein
F;1-19/Domain: signal sequence #status predicted <SIG>
F;20-104/Product: cell fusion glycoprotein F2 #status predicted <FG2>
F;105-546/Product: cell fusion glycoprotein F1 #status predicted <FG1>
F;109-133/Domain: transmembrane #status predicted <TM1>
F;485-513/Domain: transmembrane #status predicted <TM2>
F;25-57/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 30.5%; Score 74; DB 1; Length 546;
Best Local Similarity 59.3%; Pred. No. 0.12;
Matches 16; Conservative 2; Mismatches 5; Indels 4; Gaps 1;

QY 16 FEGLSIRKGVIVHRLGV----GGEHW 38
DB 281 YPSLSIRKGVIVHRLSVSYNIGSGEW 307
||| :|||||
| :|

RESULT 14
JQ2223
cell fusion protein F0 precursor - phocine distemper virus
N;Contains: F1 and F2 chains
C;Species: phocine distemper virus
C;Date: 14-Jul-1994 #sequence_revision 14-Jul-1994 #text_change 09-Jul-2004
C;Accession: JQ2223
R;Visser, I.K.G.; van der Heijden, R.W.J.; van de Bildt, M.W.G.; Kenter, M.J.H.; Oerfell,
J. Gen. Virol. 74, 1989-1994, 1993
A;Title: Fusion protein gene nucleotide sequence similarities, shared antigenic sites and
e virus entity.
A;Reference number: JQ2223; MUID:93389459; PMID:8376973
A;Accession: JQ2223
A;Molecule type: mRNA
A;Residues: 1-542 <VTS>
A;Cross-references: UNIPROT:Q7LZY1; GB:L07075
A;Note: The authors translated the codon ATC for residue 4 as Leu
C;Comment: This fusion protein F0 is cleaved into F1 and F2 chains.
C;Genetics:
A;Gene: F
C;Superfamily: parainfluenza virus cell fusion protein
C;Keywords: glycoprotein; membrane fusion; transmembrane protein
F;1-15/Domain: signal sequence #status predicted <SIG>

F;16-542/Product: fusion protein #status predicted <MAT>
 F;16-99/Product: F2 chain #status predicted <F2C>
 F;105-542/Product: F1 chain #status predicted <F1C>
 F;105-135/Region: hydrophobic
 F;486-512/Domain: transmembrane #status predicted <TMM>
 F;21,53,59,397/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 29.6%; Score 72; DB 2; Length 542;
 Best Local Similarity 62.5%; Pred. No. 0.22;
 Matches 15; Conservative 2; Mismatches 3; Indels 4; Gaps 1;

QY 19 LSEIKGIVVRLRGV---GGEHW 38
 |||:|||||||:|:|
 Db 280 LSEIKGIVVRLRGVSYNIGQEW 303

RESULT 15

S47034

cell fusion protein precursor - porpoise morbillivirus
 N/Alternate names: F protein

C/Species: porpoise morbillivirus

C/Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 09-Jul-2004

C/Accession: S47034

R/Bolt, G.; Gottschalk, E.; Blixenkron-Moeller, M.; Wlahaupt, R.G.A.; Welsch, M.J.; Es

submitted to the EMBL Data Library, July 1994

A/Description: Nucleotide sequence comparisons of the F and M genes of cetacean morbilli

A/Reference number: S47034

A/Accession: S47034

A/Molecule type: mRNA

A/Residues: 1-552 <BOL>

A/Cross-references: UNIPROT:Q66147; EMBL:X80757; NID:G520639; PIDN:CAA56731.1; PID:G5206

A/Experimental source: isolate Uster 88

A/Note: the source is designated as Cetacean morbillivirus

C/Superfamily: parainfluenza virus cell fusion protein

F;1-25/Domain: signal sequence #status predicted <SIG>

F;26-552/Product: fusion protein #status predicted <MAT>

Query Match 29.6%; Score 72; DB 2; Length 552;

Best Local Similarity 62.5%; Pred. No. 0.22;

Matches 15; Conservative 2; Mismatches 3; Indels 4; Gaps 1;

QY 19 LSEIKGIVVRLRGV---GGEHW 38
 |||:|||||||:|:|
 Db 290 LSEIKGIVVRLRGVSYNIGQEW 313

Search completed: February 8, 2005, 17:31:34
 Job time : 40 secs

